

Evaluating surface ozone- temperature relationships over the eastern U.S. in chemistry-climate models

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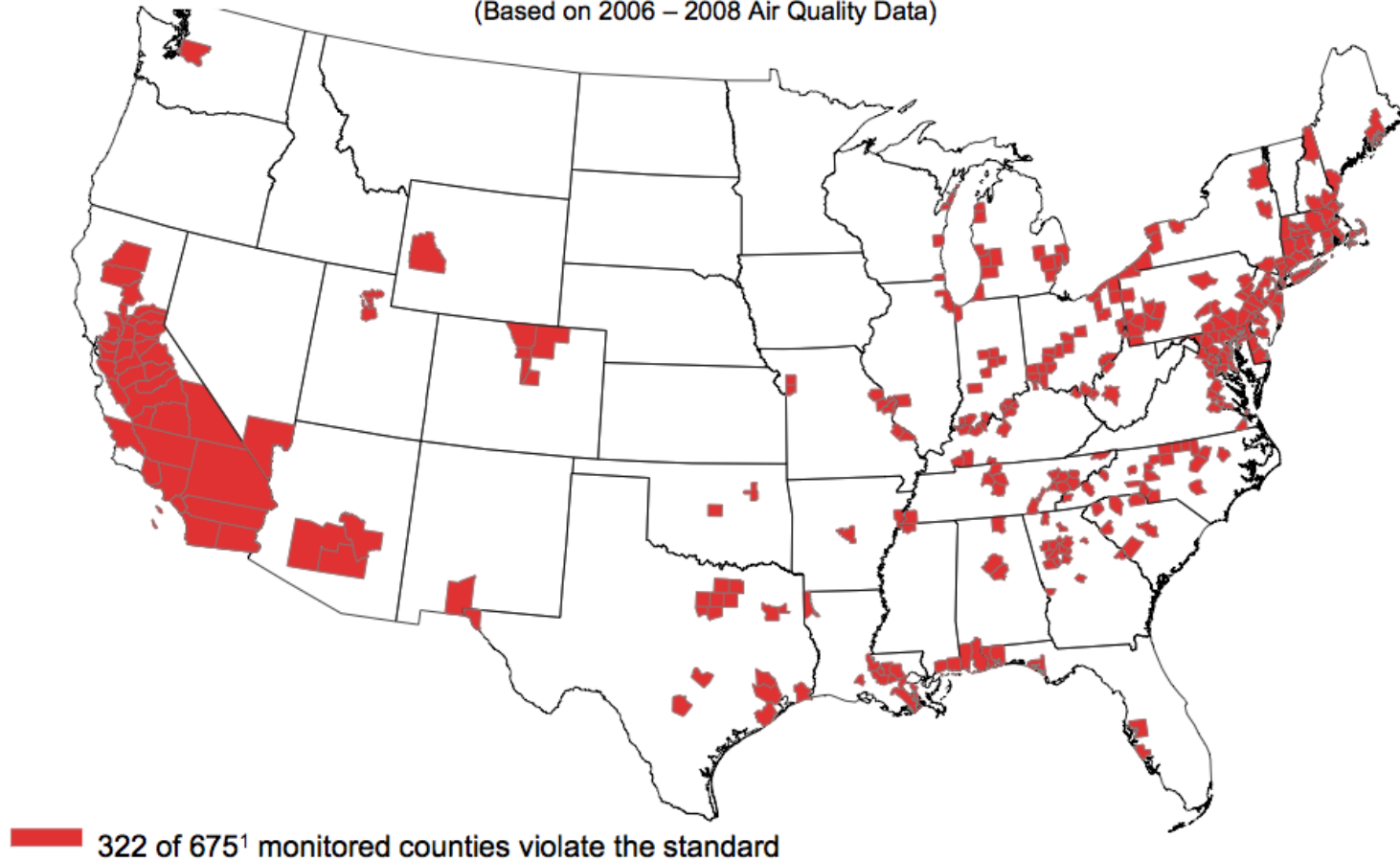


**A GFDL Lunchtime Seminar
August 17th, 2011**



The U.S. O₃ problem: over 150 million people breathe air deemed unhealthy by EPA

Counties With Monitors Violating the March 2008 Ground-Level Ozone Standards
0.075 parts per million
(Based on 2006 – 2008 Air Quality Data)



About 1 in 2 people in the U.S. live in these areas

[EPA, 2010]

(*) To attain this standard:

the 3-year average of the fourth-highest daily maximum 8-hour average ozone concentrations measured at each monitor within an area over each year must not exceed 0.075 ppm. (effective May 27, 2008)

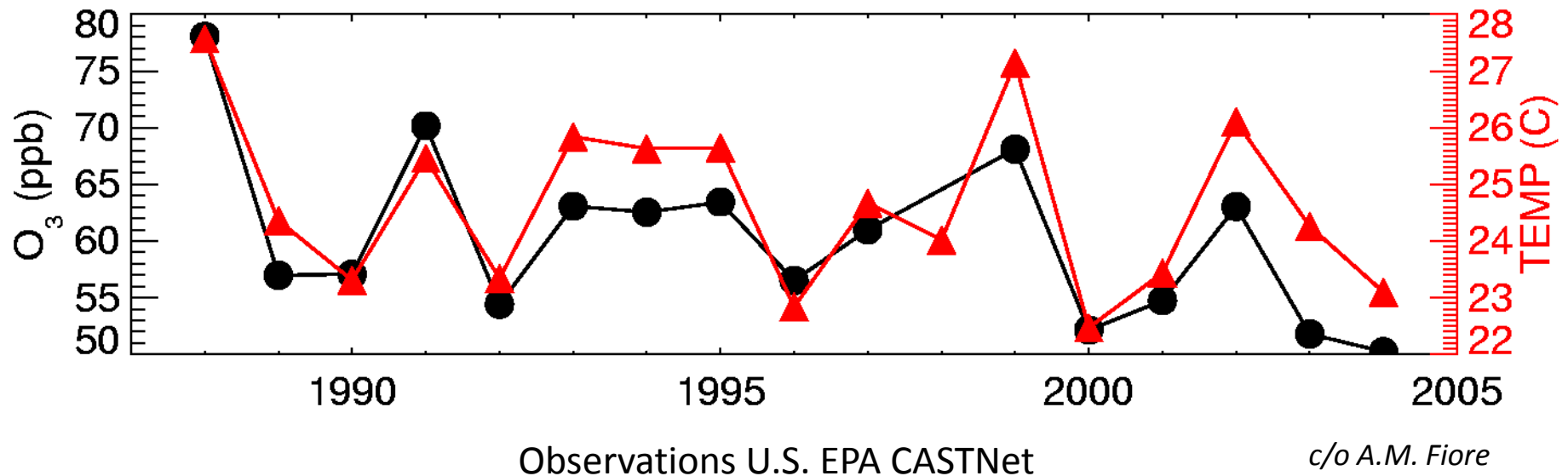
Presentation Outline

- Implications for air quality response to climate change
- Characterizing the O₃-temperature relationship over the eastern US
- How well does a chemistry-climate model represent this relationship, particularly in light of a modeled summertime O₃ bias over this region?
- Do biases in surface temperature contribute to this modeled O₃ bias?

Surface O₃ varies strongly with temperature

Observational studies have shown strong correlation between surface temperature and O₃ concentrations (Bloomer et al., 2009; Camalier et al., 2007; Cardelino and Chameides, 1990; Clark and Karl, 1982; Korsog and Wolff, 1991)

Avg. July Daily Max 8-hour O₃ and 10am-5pm Temp.
Penn State PA 41N -78E 378m



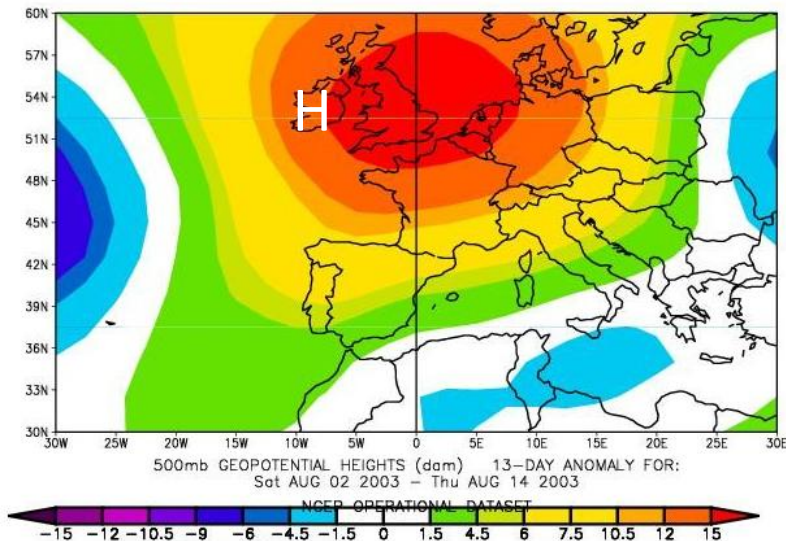
Year-to-year temperatures have been observed to be strongly correlated with ozone

Strong relationship between weather and pollution implies that changes in climate will influence air quality

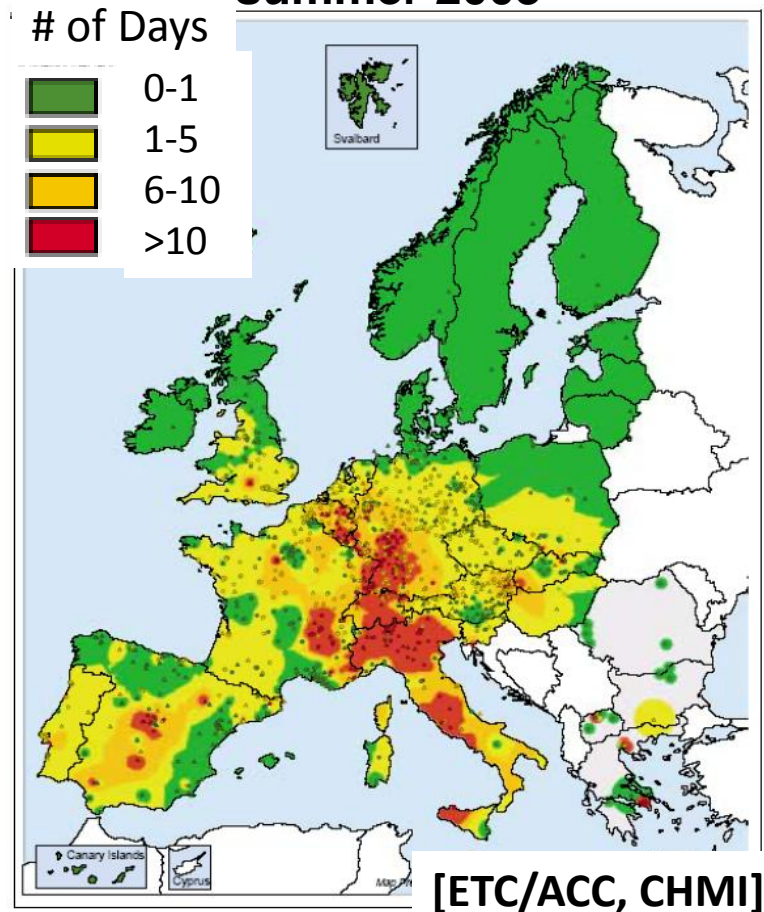
Severe O₃ pollution events associated with air stagnation in Europe

Stagnant high pressure system over Europe

(500 hPa geopotential anomaly relative to 1979-1995 for 2-14 August, NCEP)



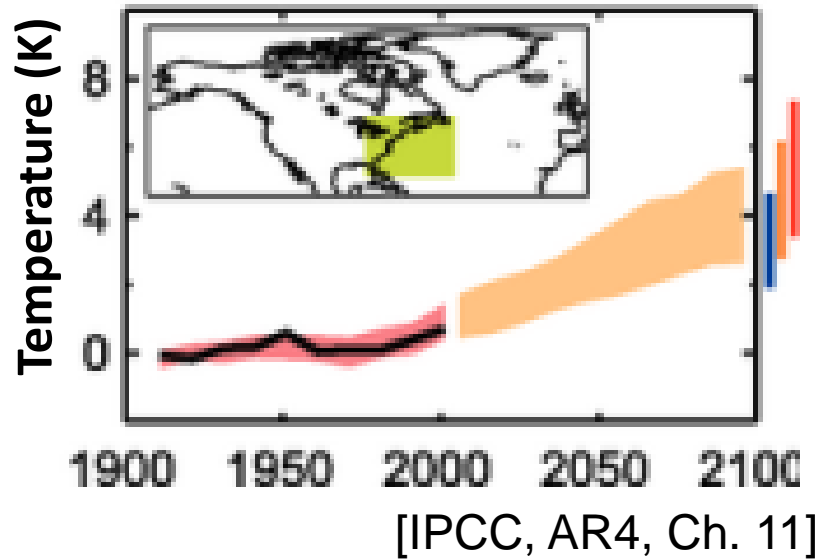
8-hr O₃ exceedances of 90 ppbv, summer 2003



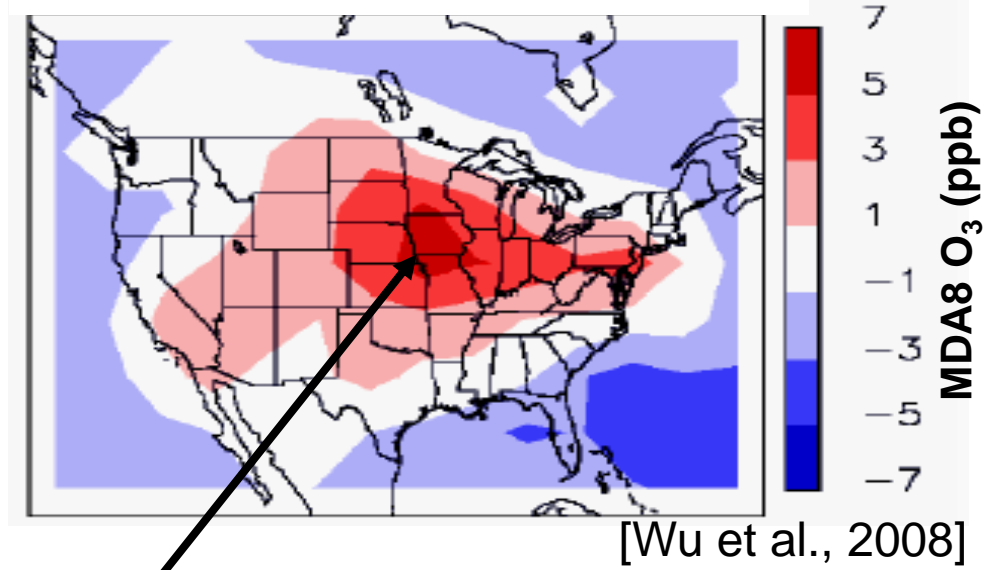
Correlation between surface O₃ and temperature is strongly associated with air stagnation

What does a warming planet imply for surface O_3 in the eastern US?

eastern US



Changes in summer 8-h avg. daily maximum ozone from 2000-2050 climate change



Higher temperatures, stagnation

- Models agree that 2000-2050 climate change will **increase surface ozone most over polluted regions (3-5 ppb)**.
- Most models find climate change will **exacerbate pollution episodes** (up to 10 ppb) due to increased stagnation and higher temperatures.
- **Models fairly robust in simulating O_3 increases in Northeast and Midwest U.S.**

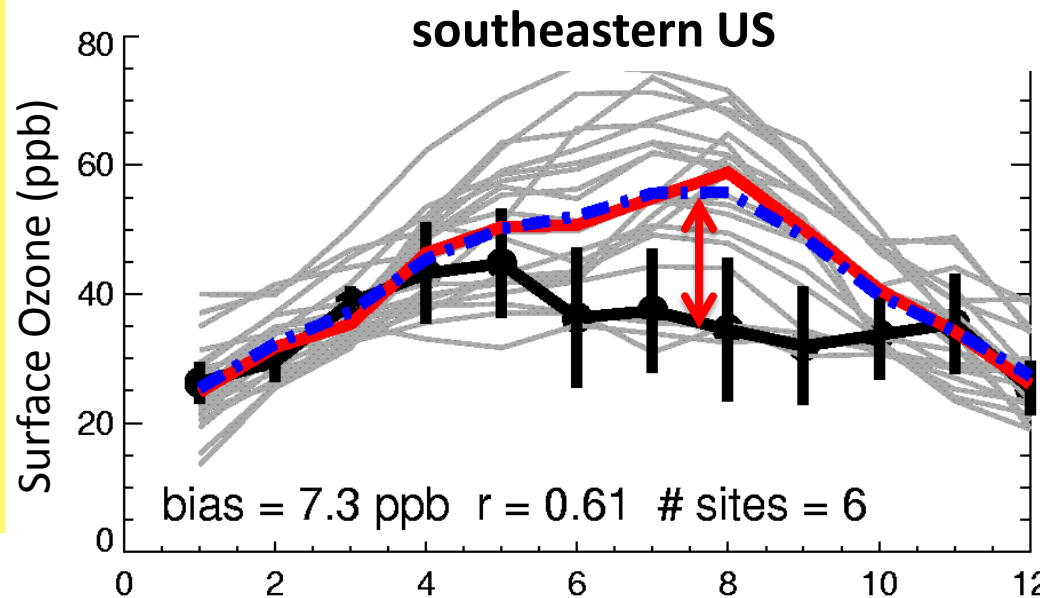
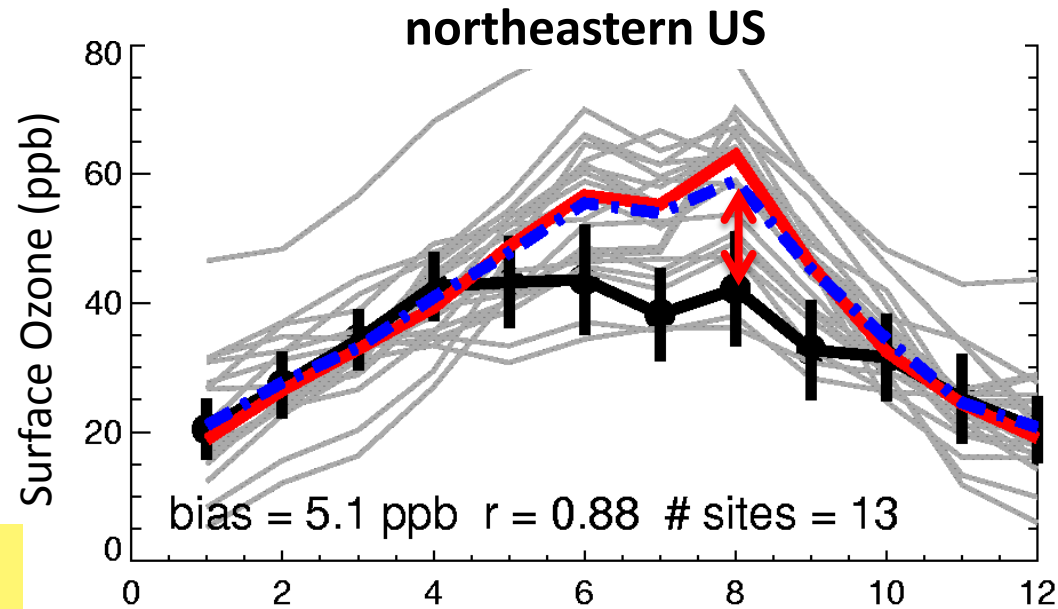
Current generation global models

How accurate are these models at reproducing observed seasonal O_3 concentrations?

Obs. = black

Ensemble mean/median = red/blue

- A pervasive **high summertime O_3 bias** exists across both eastern US regions [A.M. Fiore, L.W. Horowitz et al., 2009; Murazaki and Hess, 2006; Reidmiller et al., 2009]
- *These biases raise concern of the ability of chemistry-climate models to project accurately the response of air pollution to climate change*



Motivating Questions

- 1) Can we characterize long-term O_3 response to year-to-year temperatures for regions over the eastern US for purposes of chemistry-climate model evaluation?
- 2) Despite a known modeled ozone bias, can the GFDL AM3 chemistry-climate model represent the **response** of surface O_3 to interannual variations in temperature?
- 3) Do modeled temperature biases contribute to these modeled ozone biases?

We hypothesize that:

Adequate representation of observed, climatological O_3 -temperature relationships will help to build confidence in future projections of the air quality response to changes in climate

Representing the O₃ – temperature relationship

We use temperature as a proxy to synthesize the complex effects of meteorological and chemical factors influencing O₃ concentrations

We represent these aggregate effects as a total derivative, $\frac{d[O_3]}{dT}$

The O₃-temperature relationship is thought to represent at least three components in the eastern US:

$$\frac{d[O_3]}{dT} = \underbrace{\frac{\partial[O_3]}{\partial[\text{stagnation}]} \frac{\partial[\text{stagnation}]}{\partial T}}_{(1)} + \underbrace{\frac{\partial[O_3]}{\partial[\text{PAN}]} \frac{\partial[\text{PAN}]}{\partial T}}_{(2)} + \underbrace{\frac{\partial[O_3]}{\partial[\text{isoprene}]} \frac{\partial[\text{isoprene}]}{\partial T}}_{(3)} + \dots$$

[Jacob et al., 1993; Olszyna et al., 1997] *[Sillman and Samson, 1995]* *[Meleux et al., 2007; Guenther et al., 2006]*

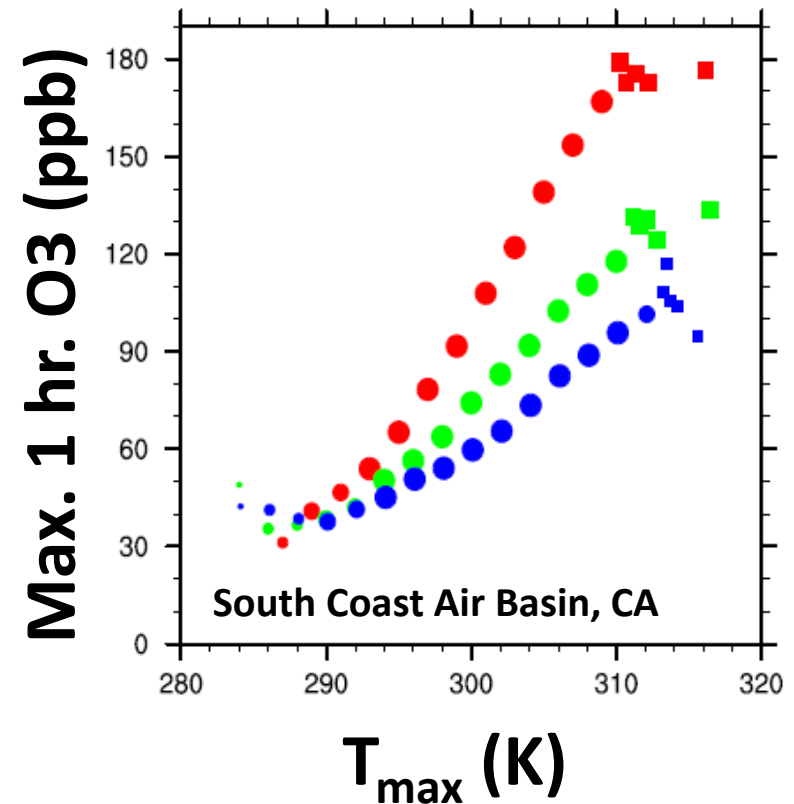
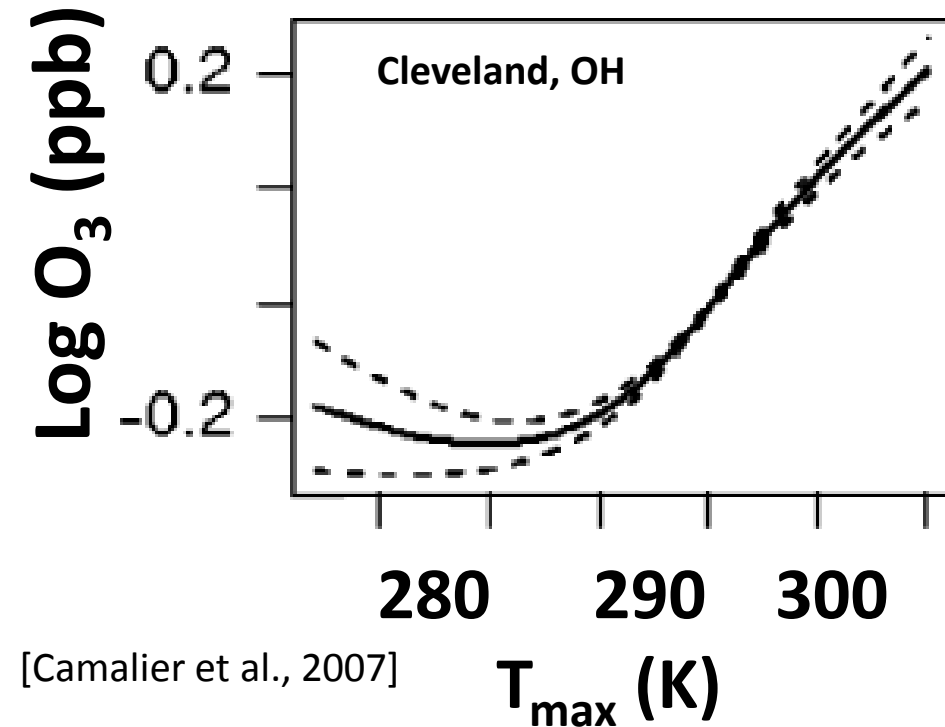
meteorology

chemistry

emissions

Over what range of temperatures does O₃ increase?

Ozone's relationship with temperature is non-linear, but it has been observed to be linear for ranges of temperatures...



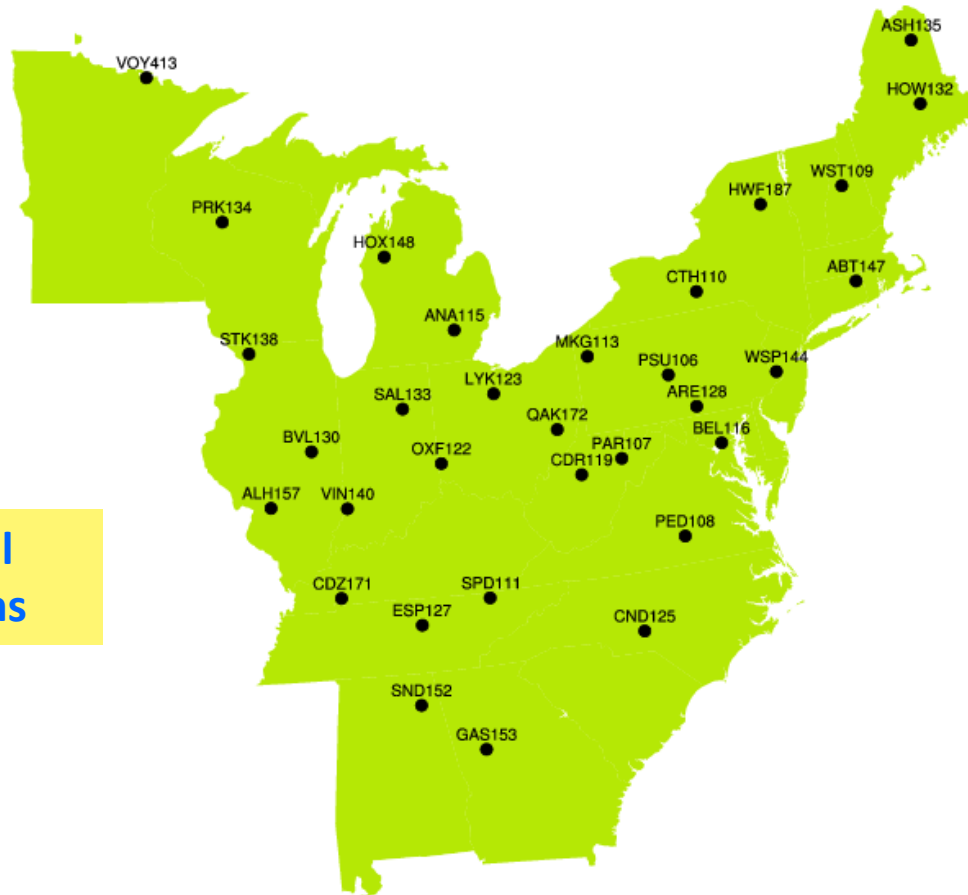
[Steiner et al., 2010]

O₃ found to typically increase linearly between 290-305 K

[Bloomer et al., 2009; Camalier et al., 2007; Sillman and Samson, 1995; Steiner et al., 2010]

Observation network

EPA CASTNet sites used and locations:



co-located meteorological
and chemical observations

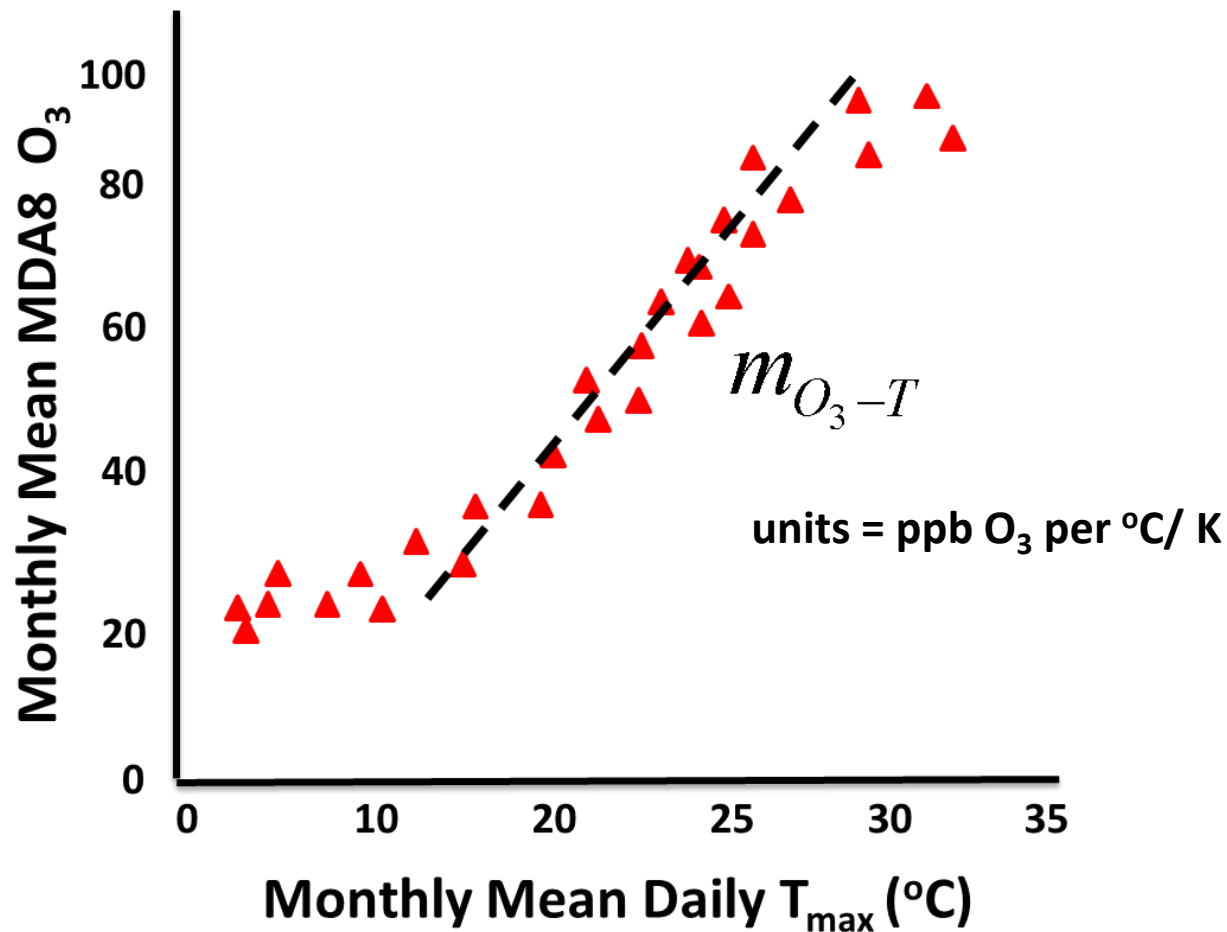
**Observations: EPA Clean Air Status and Trends Network (CASTNet;
designed for measurements to be representative of the regional scale)**

Most sites have record lengths spanning 1988-2009

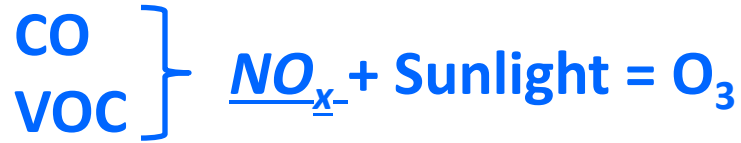
statistical approach to the O₃-temp relationship

We focus on the warm season over the eastern US where O₃ and temperature are well correlated [Dawson et al., 2007; Lin et al., 2001; Sillman and Samson, 1995]

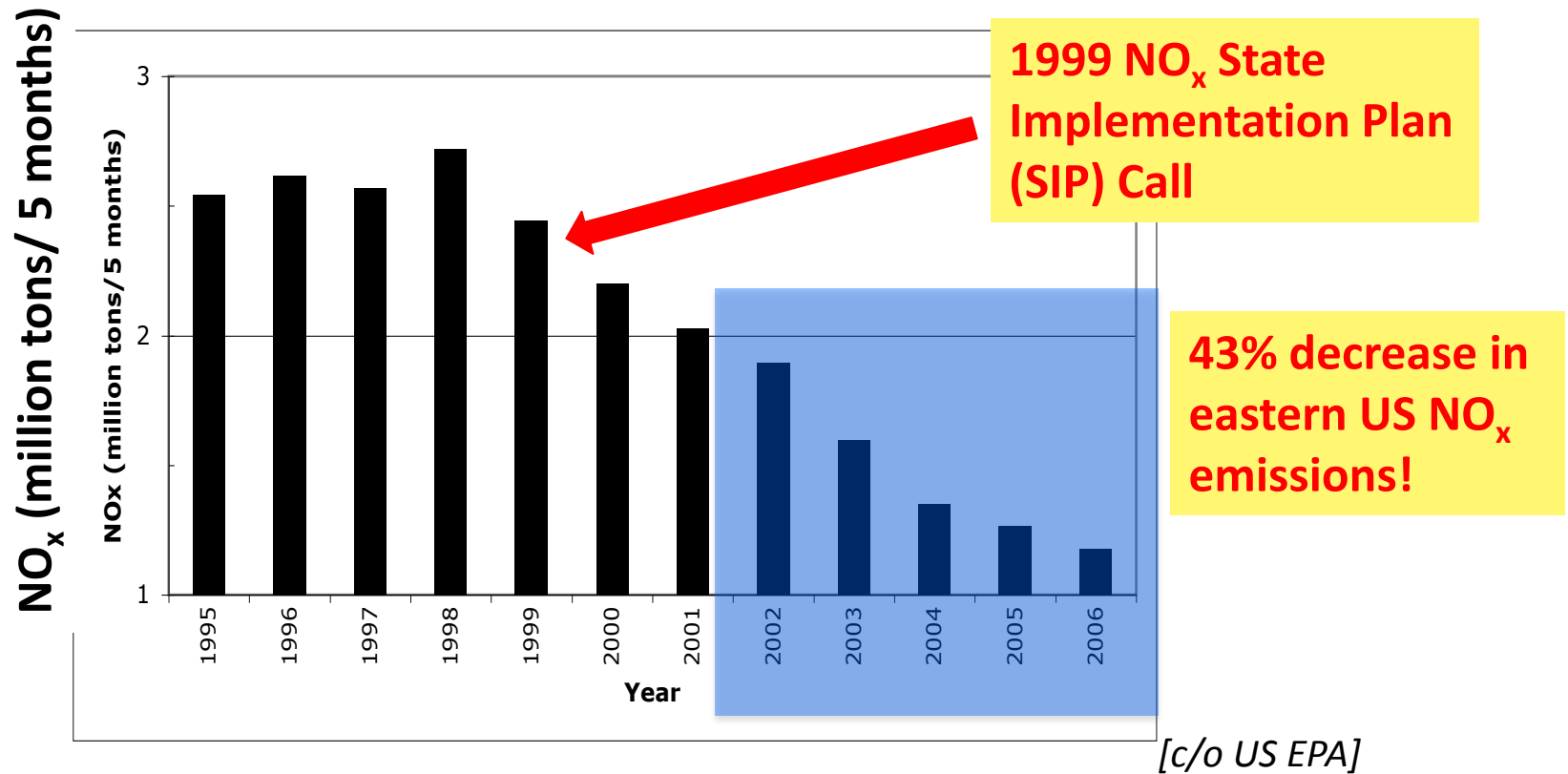
We use maximum daily 8-hour average (MDA8) O₃ and maximum daily surface temperature (T_{max}) to focus on the daytime (deep, well-mixed, boundary layer)



Humans can influence O_3 sensitivity to temperature by changing the O_3 production chemistry



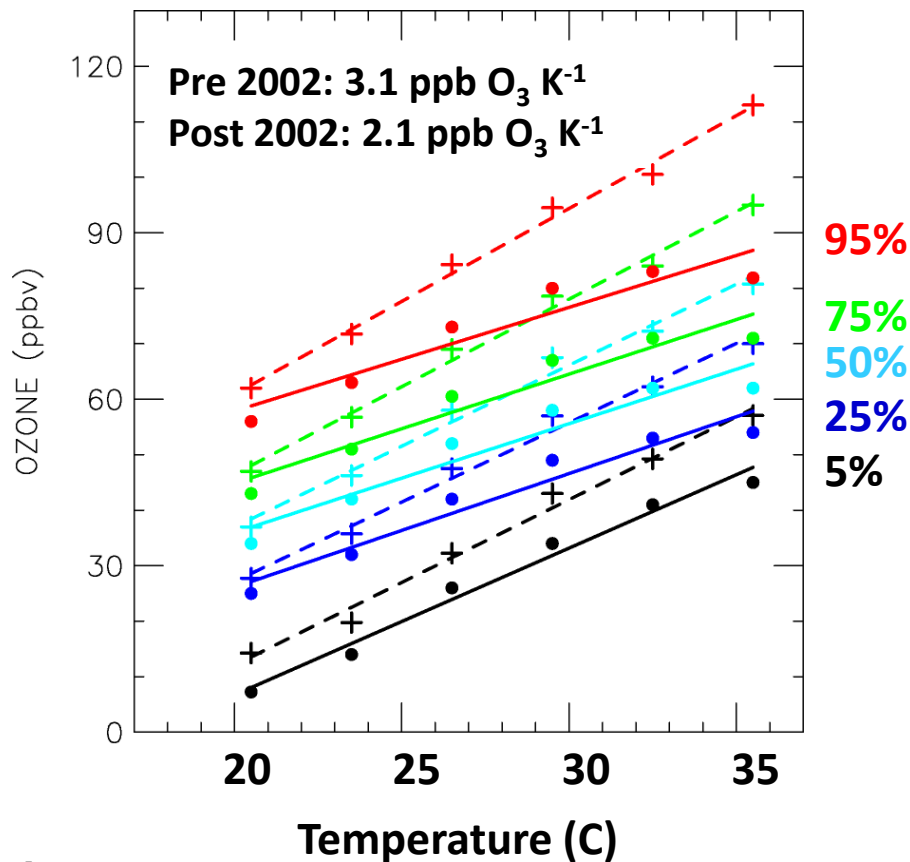
National Ozone Season NO_x Emissions from Power Plants



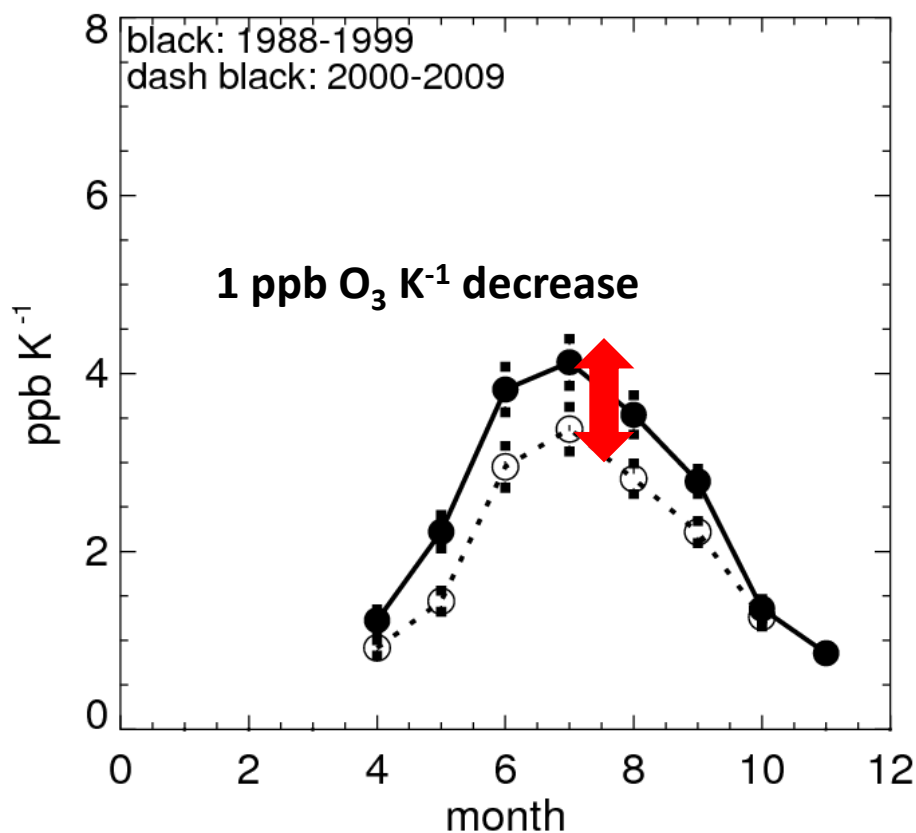
Changes to anthropogenic NO_x emissions have a substantial effect on the sensitivity of O_3 to temperature

effects of NO_x emission controls on O₃ sensitivity

Great Lakes (May-October)



Great Lakes



[Bloomer et al., 2009]

- Δ NO_x emissions decreased ozone levels over the entire distribution
- The ozone sensitivities to temperature also decreased across the distribution
- Different statistical methods reveal the same 1 ppb O₃ K⁻¹ decrease during O₃ season

AM3 simulations

- We use a 20-year (1981—2000) run of the GFDL AM3 CCM [Donner et al., 2011; Naik et al., in prep]
 - Interactive isoprene (responds to solar radiation and temperature) (MEGAN) following that used in MOZART-4 [Emmons et al., 2010]
- Forcings:
 - Observed SSTs and sea ice [Rayner et al., 2003]
 - ACCMIP aerosol and O3 precursor emissions (as used in CM3 CMIP5 simulations) [Lamarque et al., 2010]

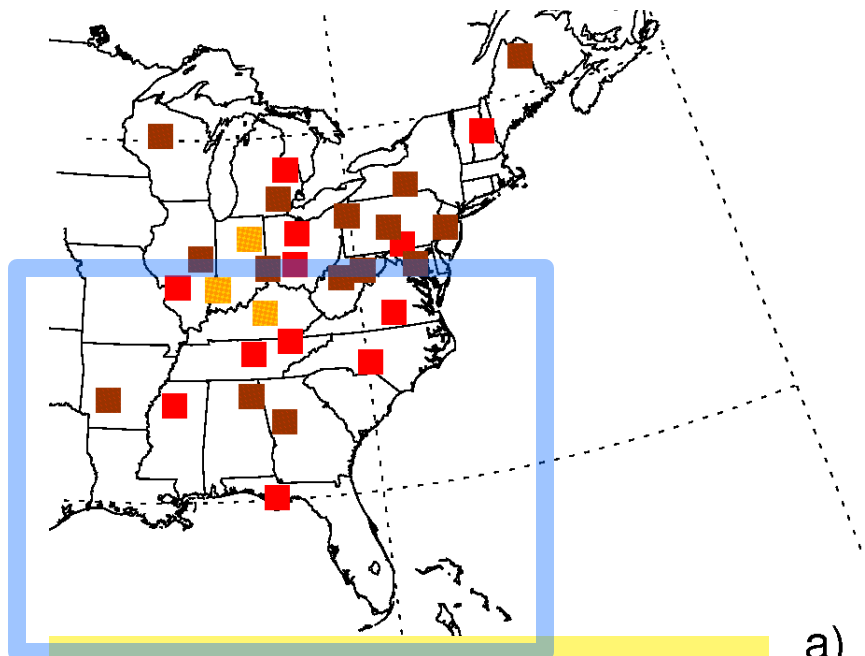
Little to no change in eastern US NO_x emissions in model run

	1980	1990	2000
NO _x	4.23 Tg N	4.40 Tg N	4.29 Tg N

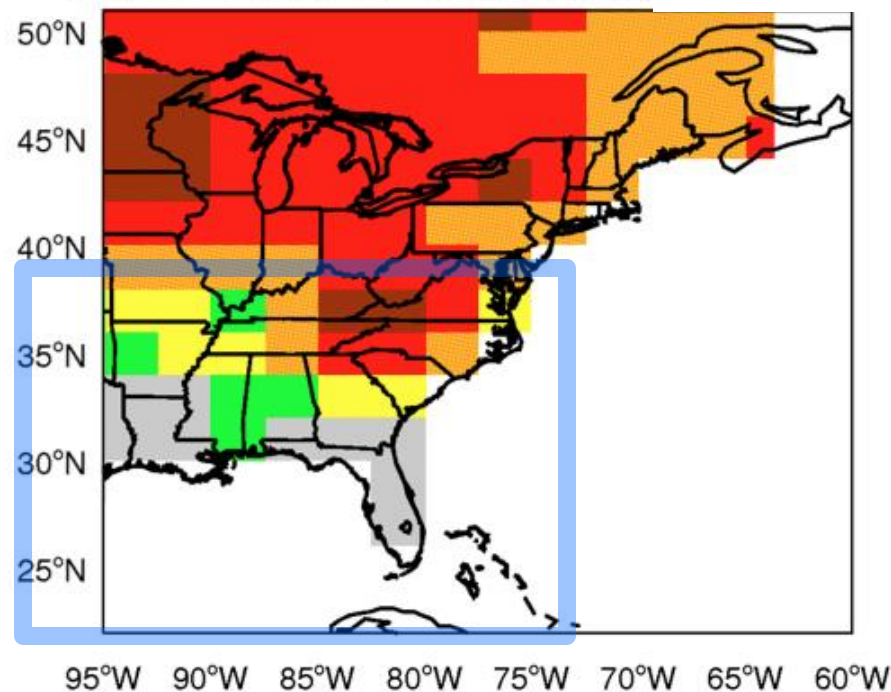
“climatological” O₃-temperature relationships

correlating MDA8 O₃ and daily T_{max}

July CASTNet correlation (r) 1988-1999



July GFDL AM3 correlation (r) 1981-2000



July O₃ is well correlated with T_{max}

a)

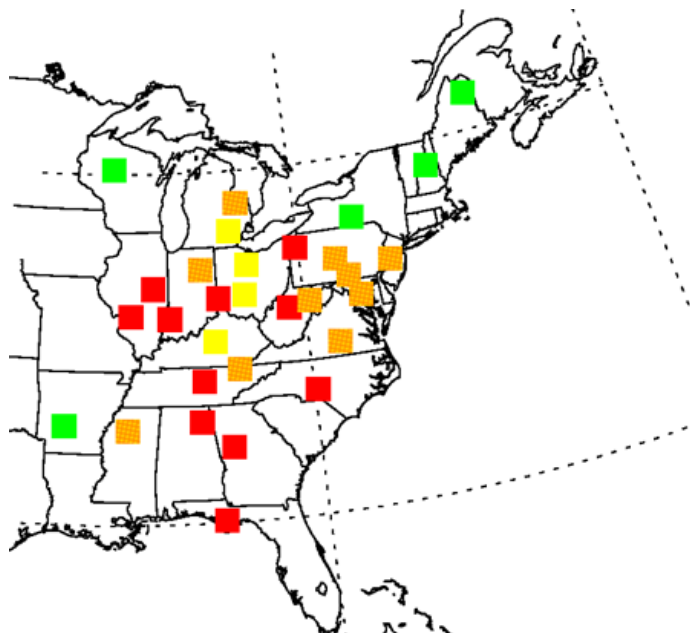


correlations too low in southern US

-problem representing inflow of marine air, convective ventilation, isoprene chemistry in this region?

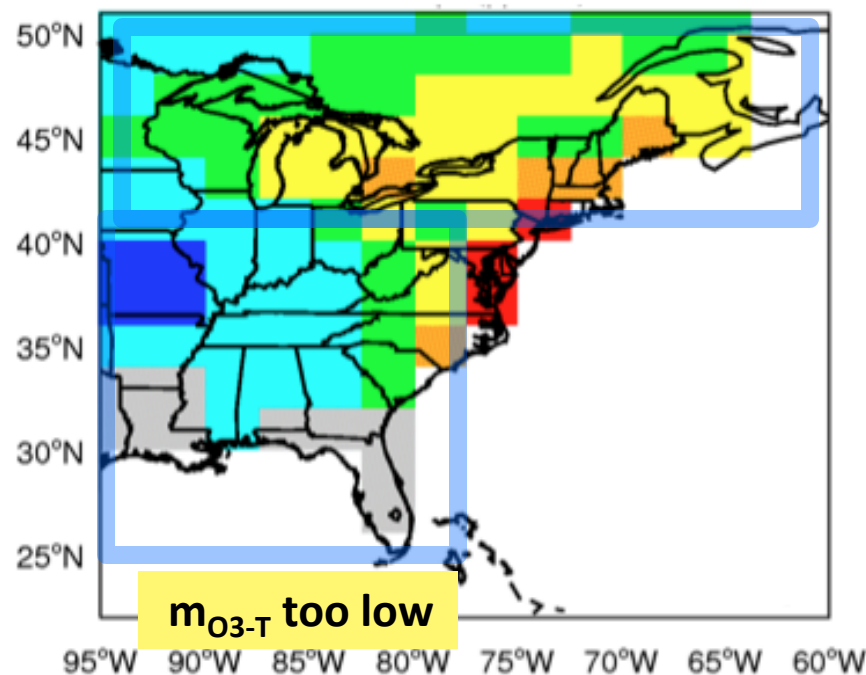
“climatological” O₃-temperature relationships

July CASTNet slope (ppb K⁻¹) 1988-1999



July m_{O_3-T} highest in Mid-Atlantic and southern US

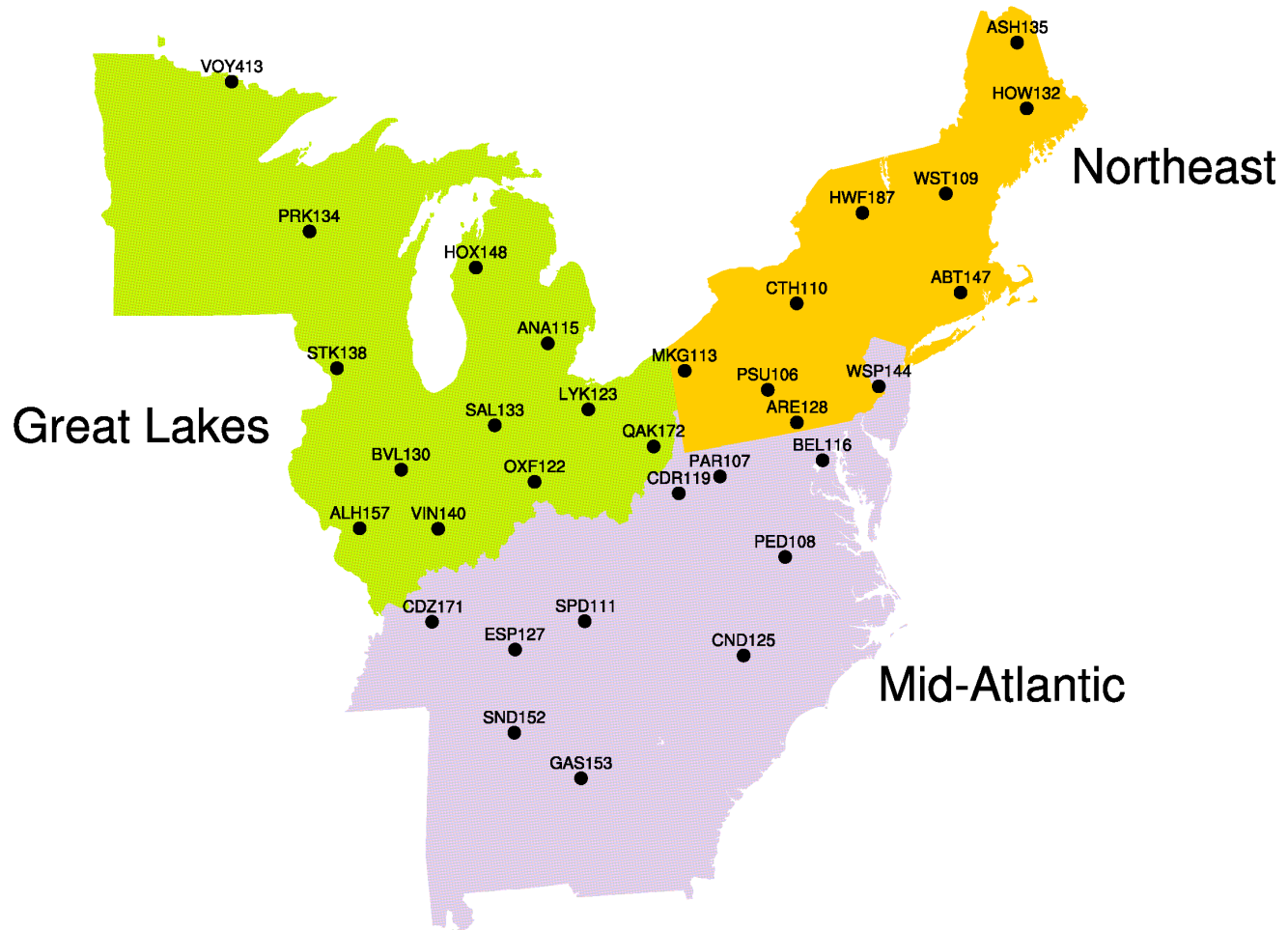
July CASTNet slope (ppb K⁻¹) 1981-2000



GFDL AM3 produces range of m_{O_3-T} over Northeast and portions of the Great Lakes

Remains unclear why GFDL AM3 struggles in Mid-Atlantic and southern US

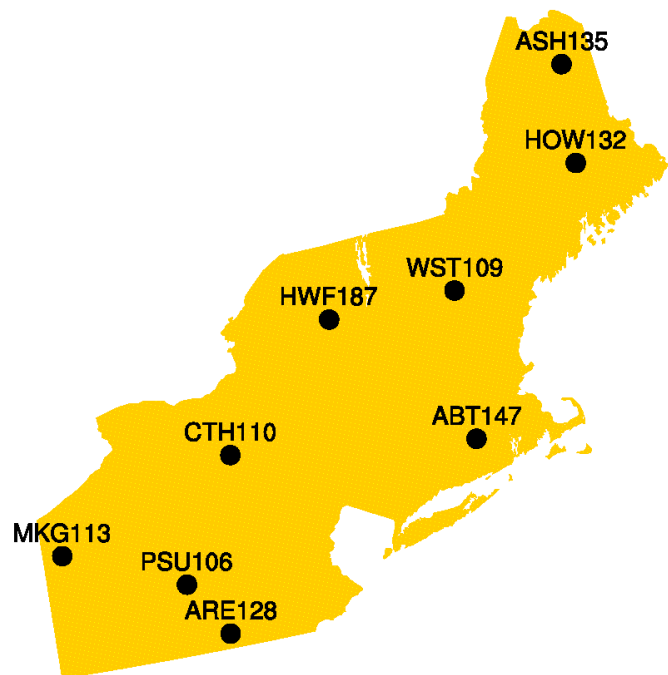
Regional approach to characterizing O₃-temperature relationships



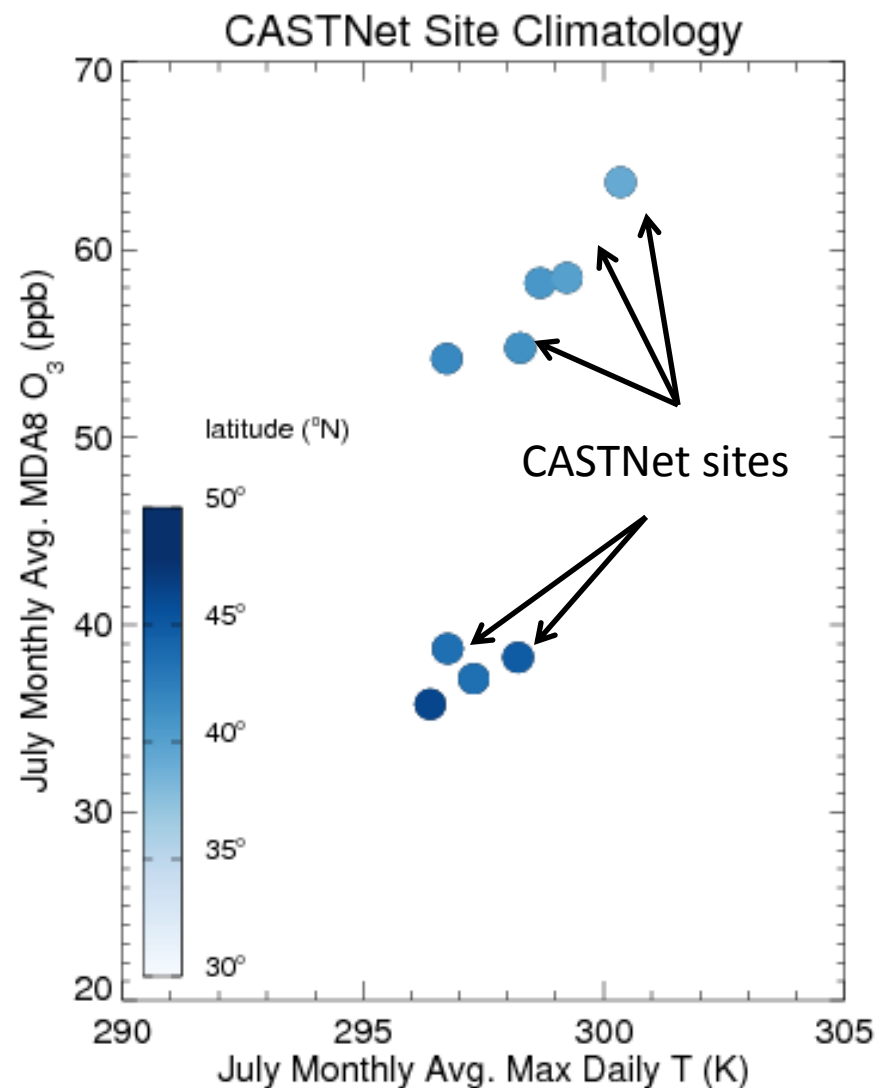
We group CASTNet sites into chemically and meteorologically coherent regions

Motivated by past statistical analyses [Bloomer et al., 2009; Eder et al., 1993; Lehman et al., 2004]

Regional approach: Northeast

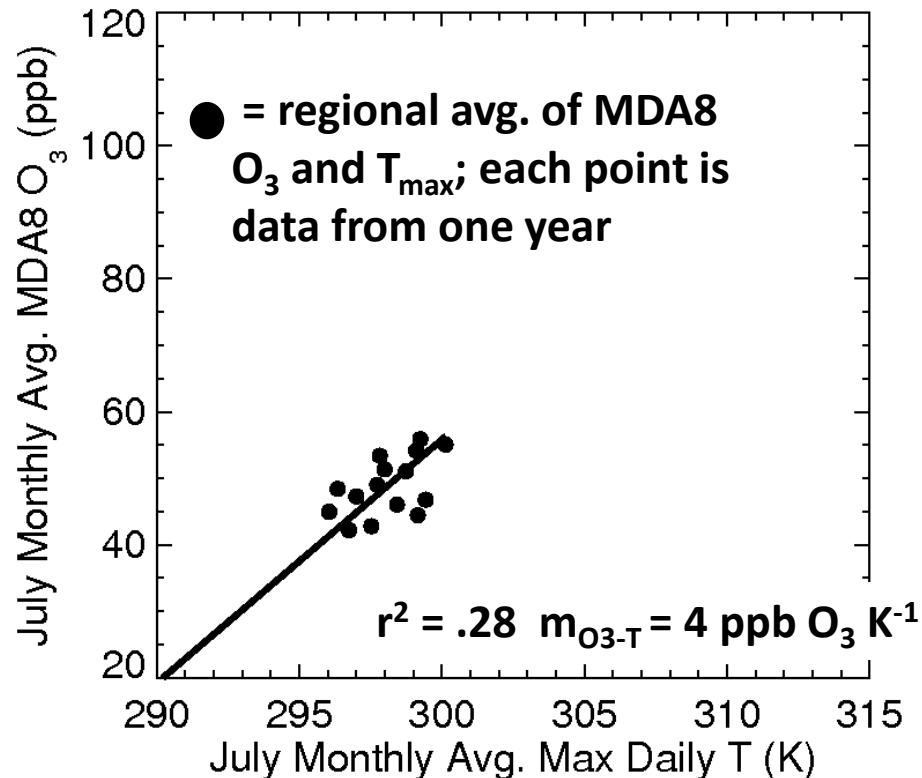


We average over all CASTNet sites to isolate the regional response of O_3 to temperature



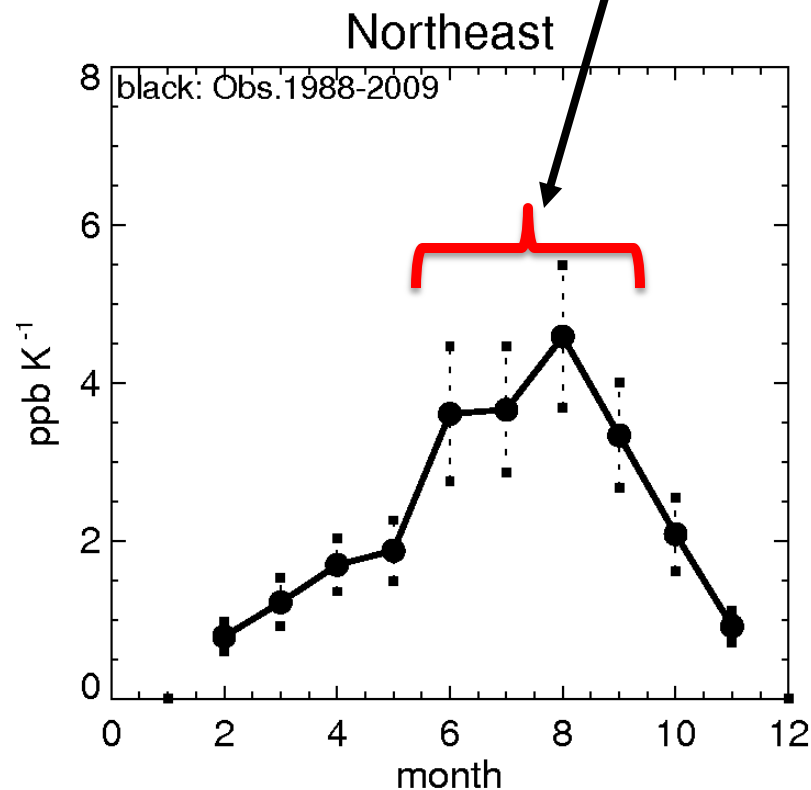
Regional approach: Northeast

Northeast

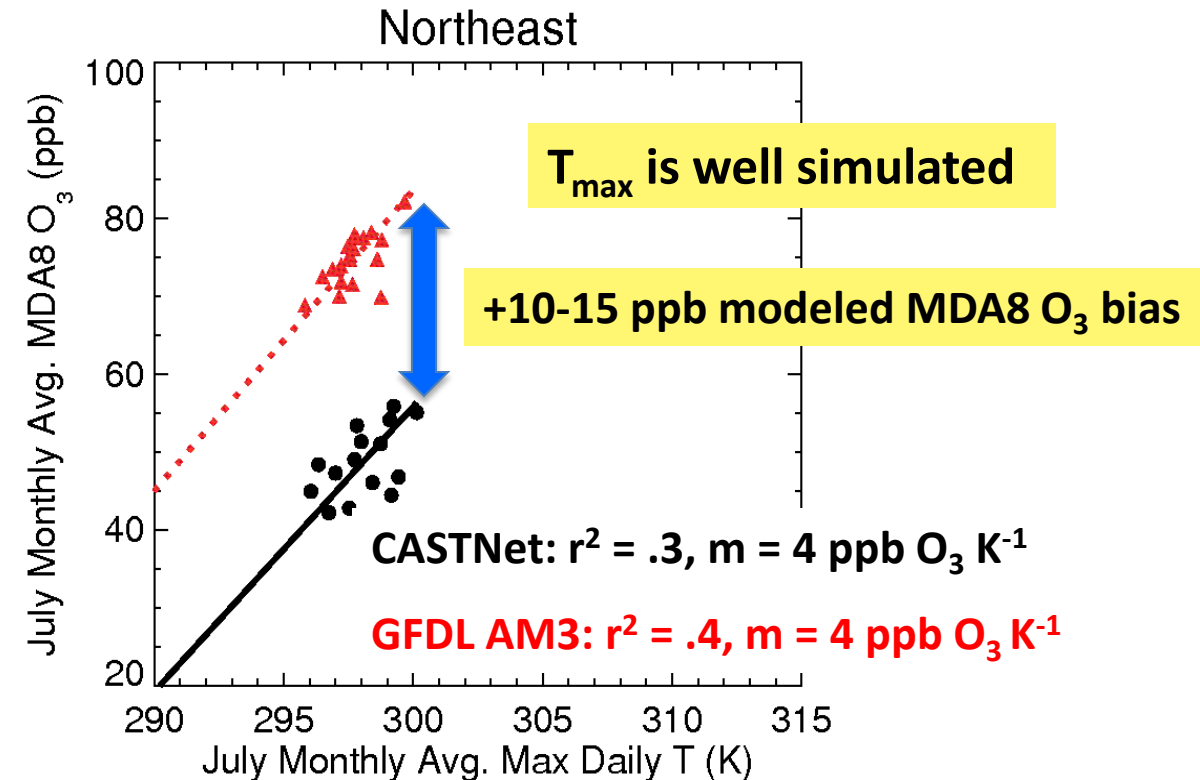


Large standard errors highlight the need for longer datasets

highest O_3 sensitivities to temperature in summer (4-5 ppb $O_3 \text{ K}^{-1}$)

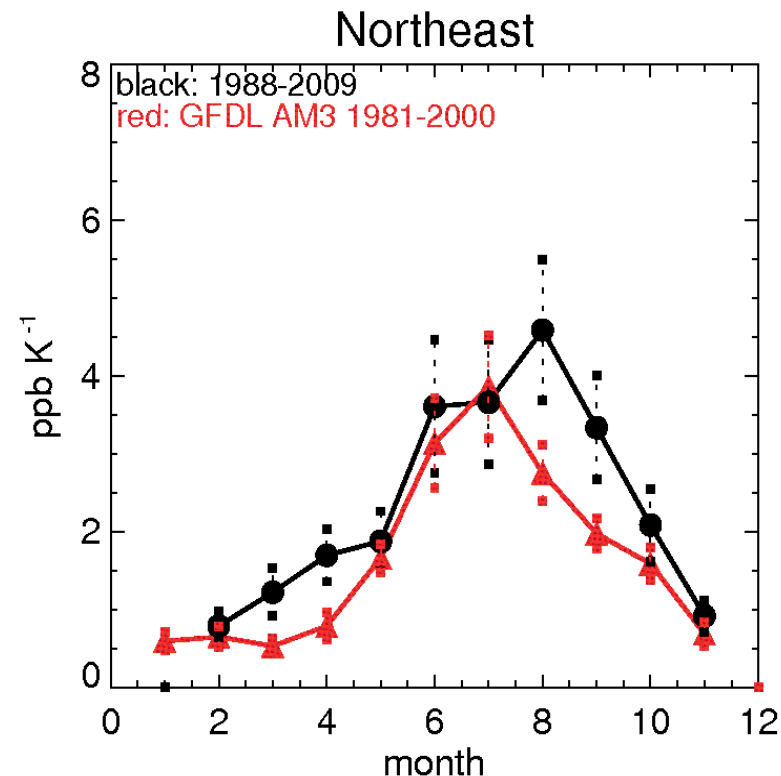


Regional approach: Northeast

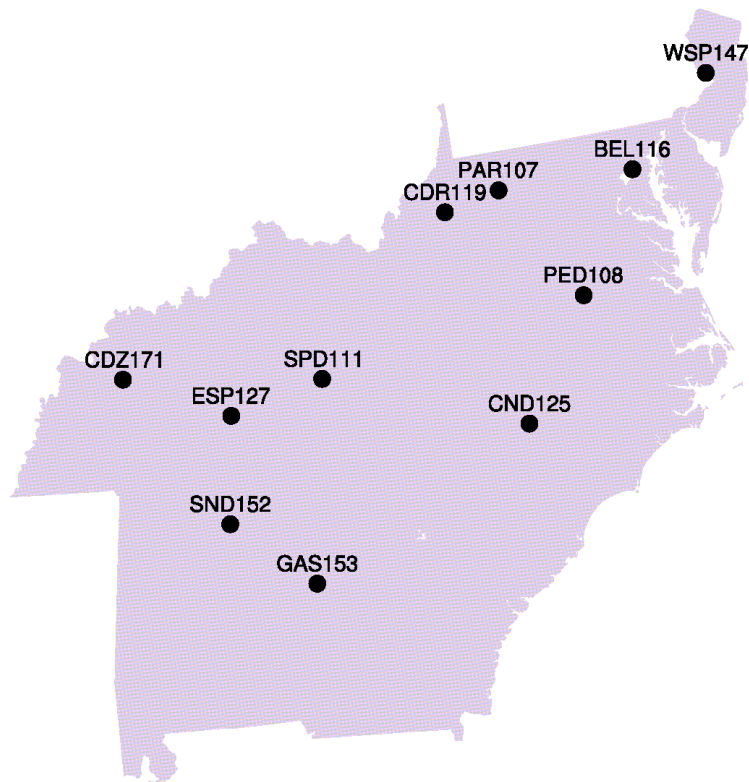


Despite the presence of excess modeled MDA8 O₃...

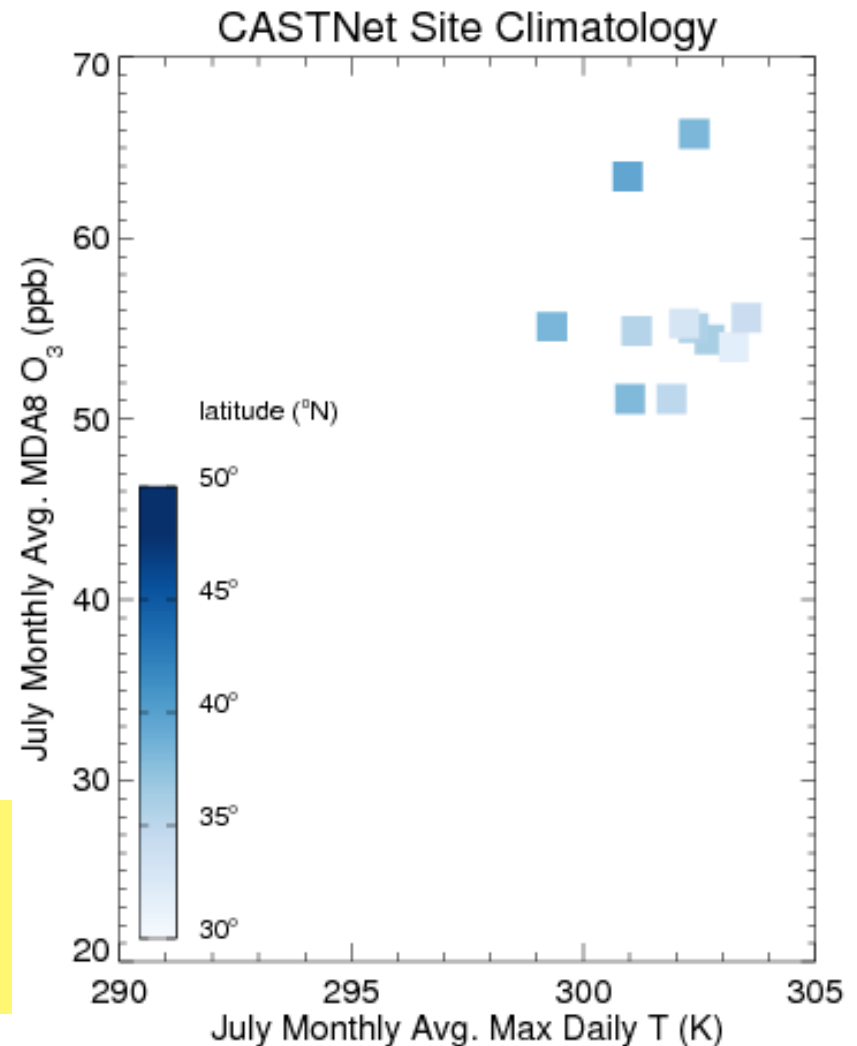
MDA8 O₃ sensitivity to year-to-year variations in T_{max} are reproduced by GFDL AM3 in the Northeast



Regional approach: Mid-Atlantic

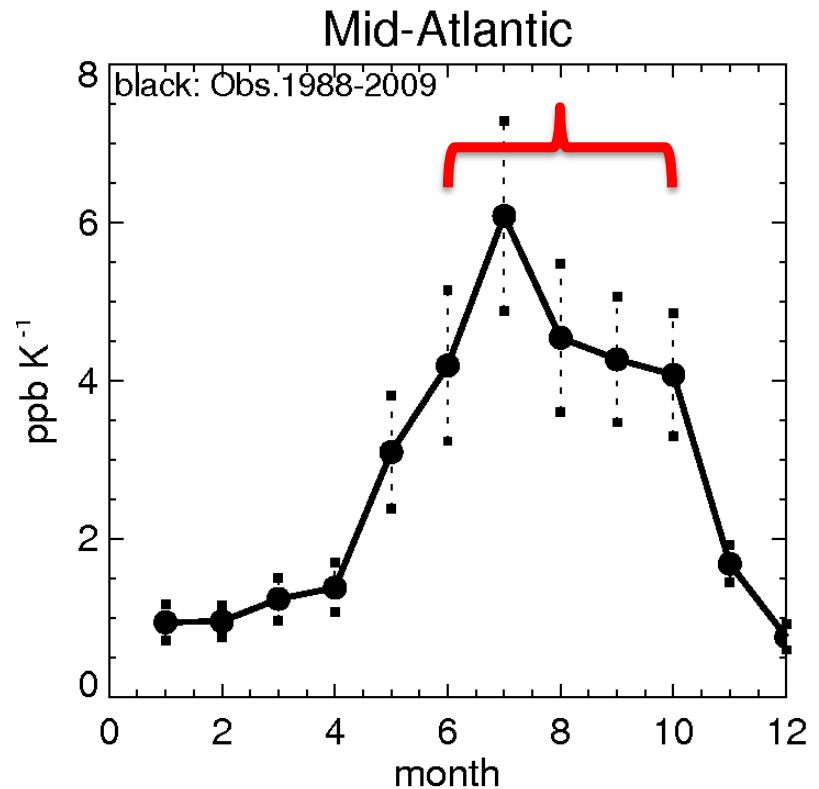
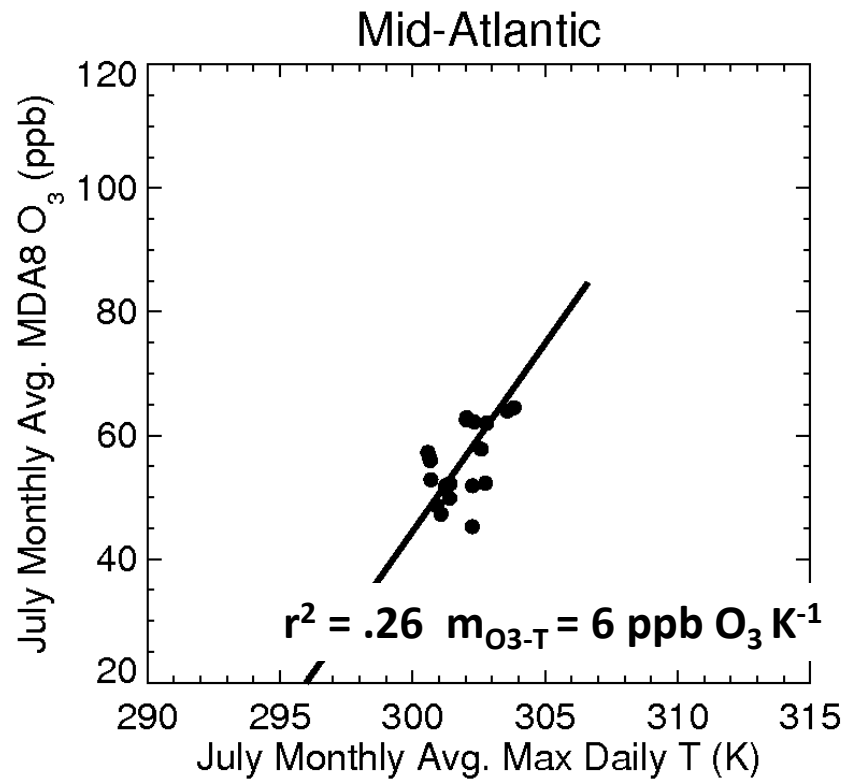


Again, we average over all CASTNet sites to isolate the regional response of O_3 to temperature

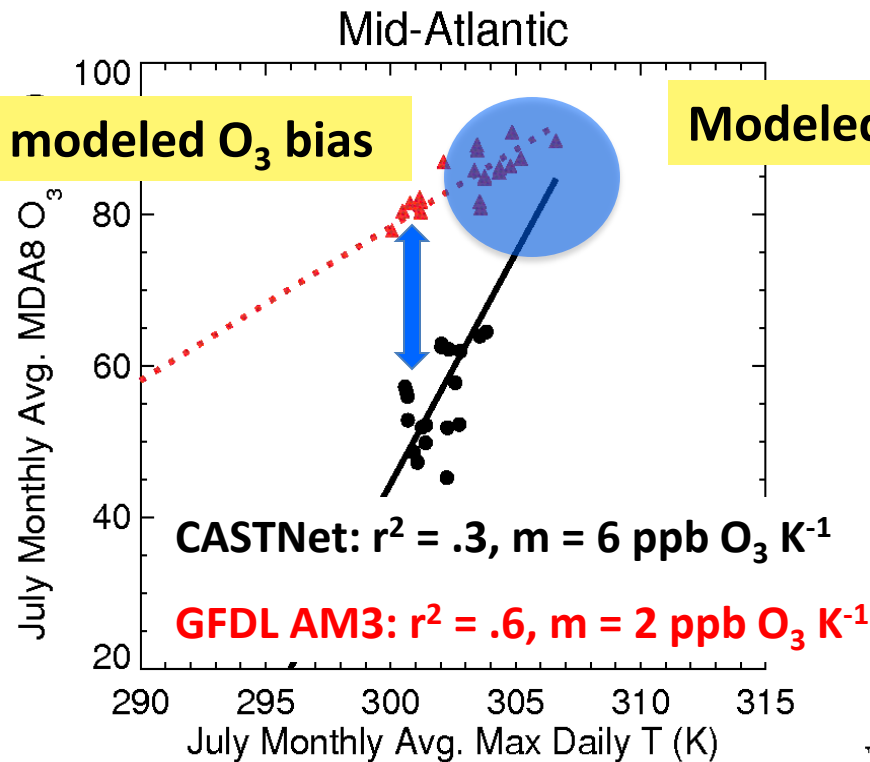


Regional approach: Mid-Atlantic

Highest O₃ sensitivities to temperature are in summer (4-6 ppb O₃ K⁻¹)

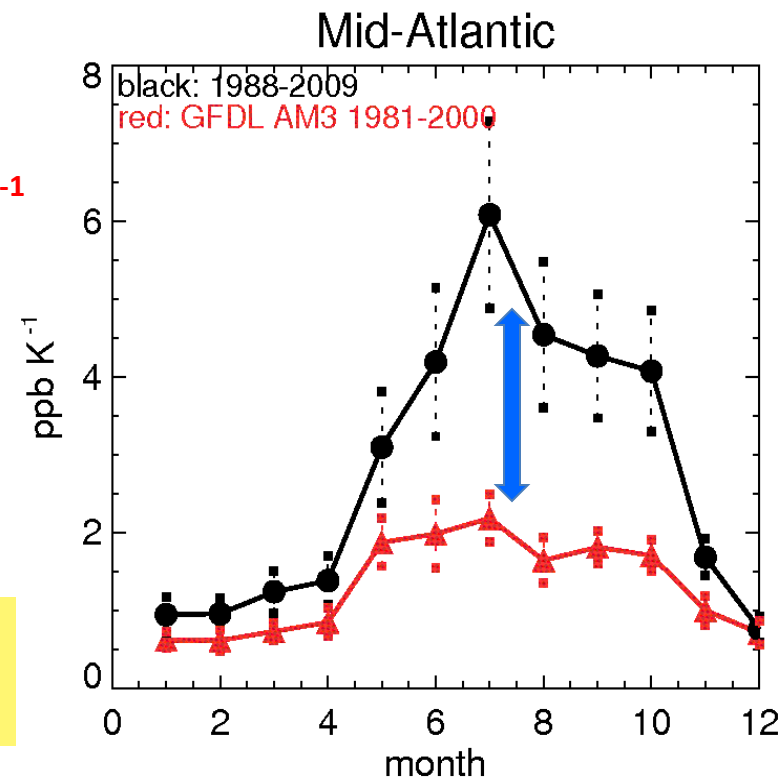


Regional approach: Mid-Atlantic



model slopes are $\sim 4 \text{ ppb } \text{O}_3 \text{ K}^{-1}$ too low in the summer months

What effect does T_{max} bias have on O_3 over the eastern US?



Modeled t_{\max} bias and O_3 sensitivity

multiply T_{\max} biases by mean regional sensitivity of 3 ppb O_3 K^{-1} $\Delta T_{\max} * 3 \left(\frac{\text{ppb } O_3}{\text{degree K}} \right)$

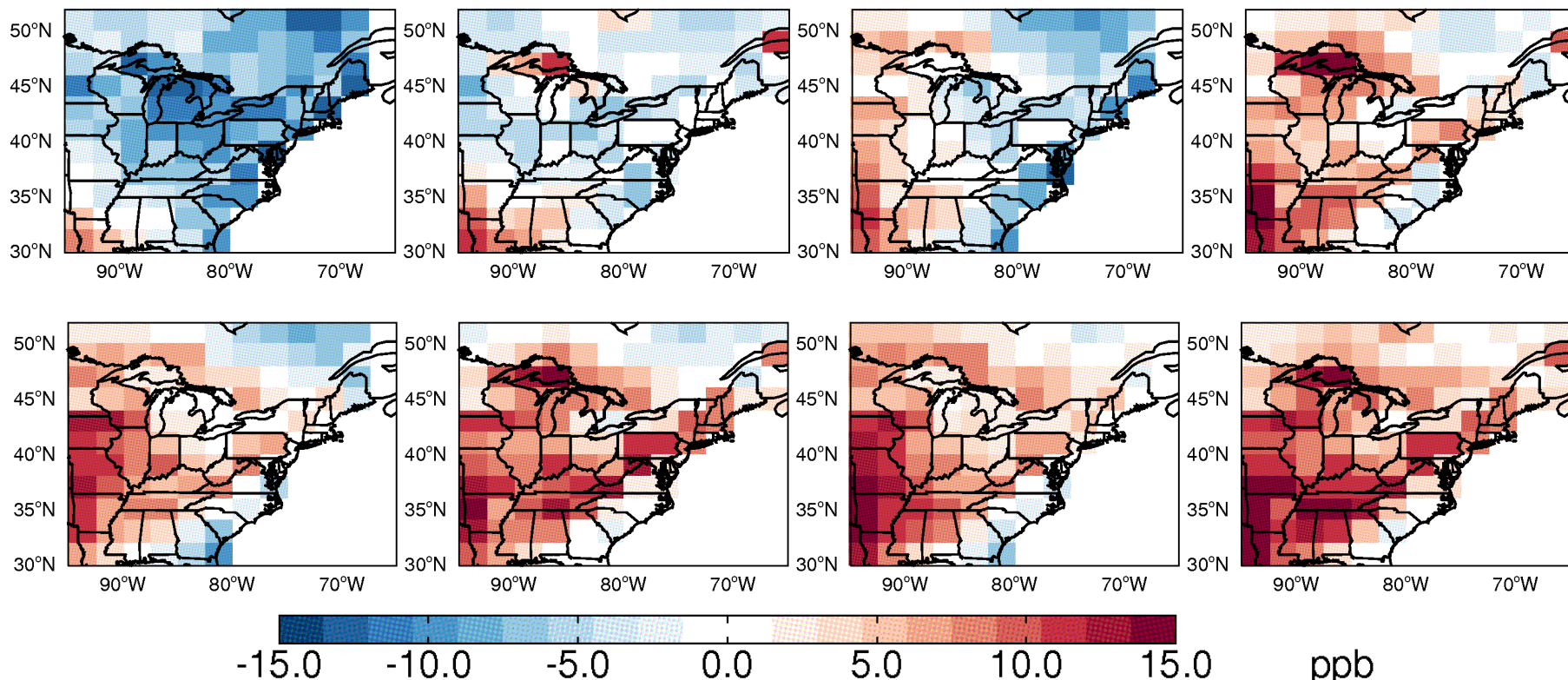
Maurer et al. [2002]

JUN

JUL

AUG

SEP



T_{\max} biases are responsible for **up to 10-15 ppb** of the summer O_3 bias, specifically in the interior of the Mid-Atlantic region

➤ cannot be the major driver of the large-scale O_3 bias

Conclusions

Despite modeled O_3 biases, the GFDL AM3 reproduces m_{O_3-T} in the Northeast, although it underestimates m_{O_3-T} by 2–4 ppb K^{-1} in the summer over the Mid-Atlantic

➤ model skill at simulating fundamental meteorological processes may be contributing to the inaccuracies in producing m_{O_3-T} over the Mid-Atlantic

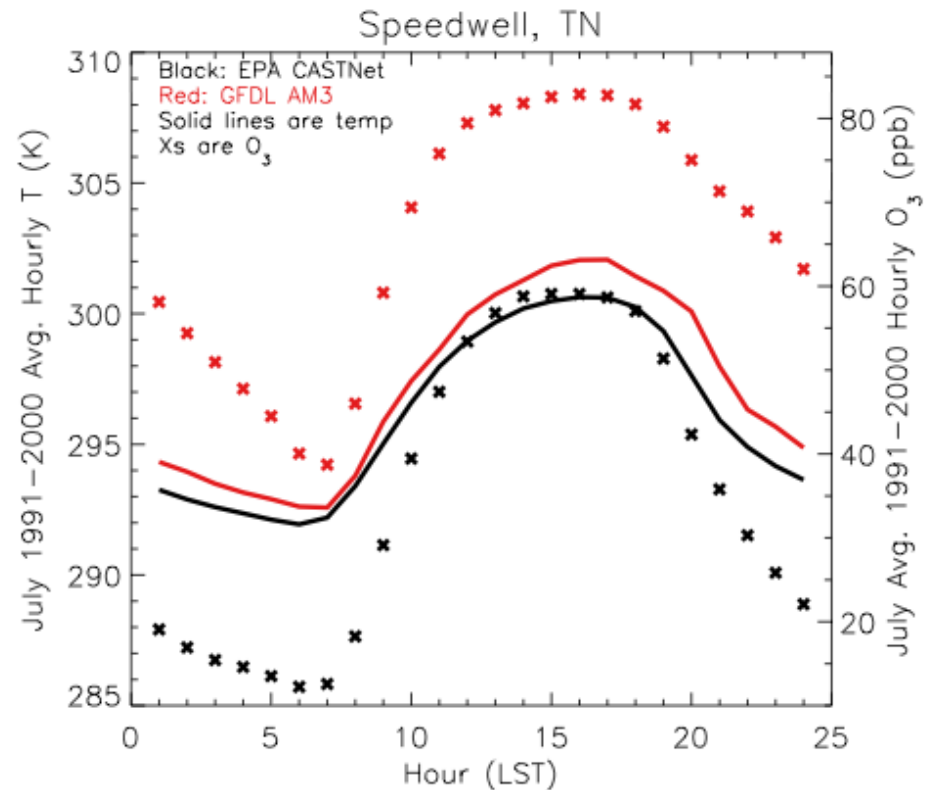
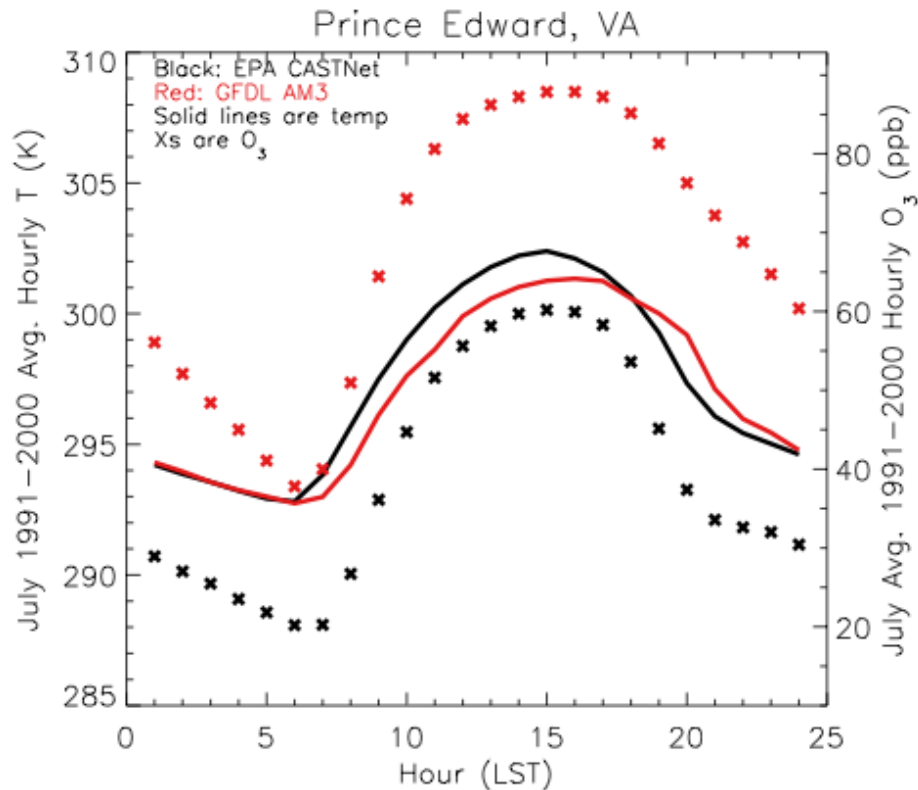
We estimate a maximum contribution of 10–15 ppb MDA8 O_3 from simulated T_{max} biases over the Mid-Atlantic

➤ We find modeled (+) T_{max} biases are not the major driver of the large-scale excess modeled O_3

supplemental slides

evaluating modeled diurnal temperature and O₃

When diurnal temperature biases are highest, are the O₃ biases correspondingly highest?

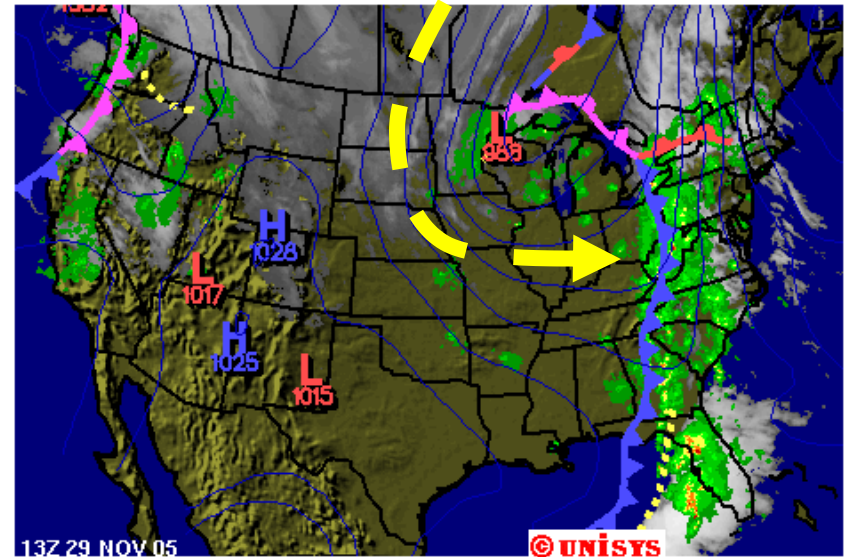


Why does the model struggle in the southeastern US?

Fundamentally different meteorological processes modulate O_3 levels in the southern and northern halves of the eastern US:

- In the Northeast: pollutant ventilation is known to be driven by migratory cyclones associated with cold fronts [Leibensperger et al., 2008; Logan, 1989, Vukovich, 1995]
- In the Southeast: Deep convection and inflow from the Gulf of Mexico are known to ventilate O_3 in the boundary layer [Li et al., 2005]
 - The skill of the GFDL AM3 chemistry-climate model in simulating these meteorological processes may help explain why O_3 -temp sensitivities are not accurately produced

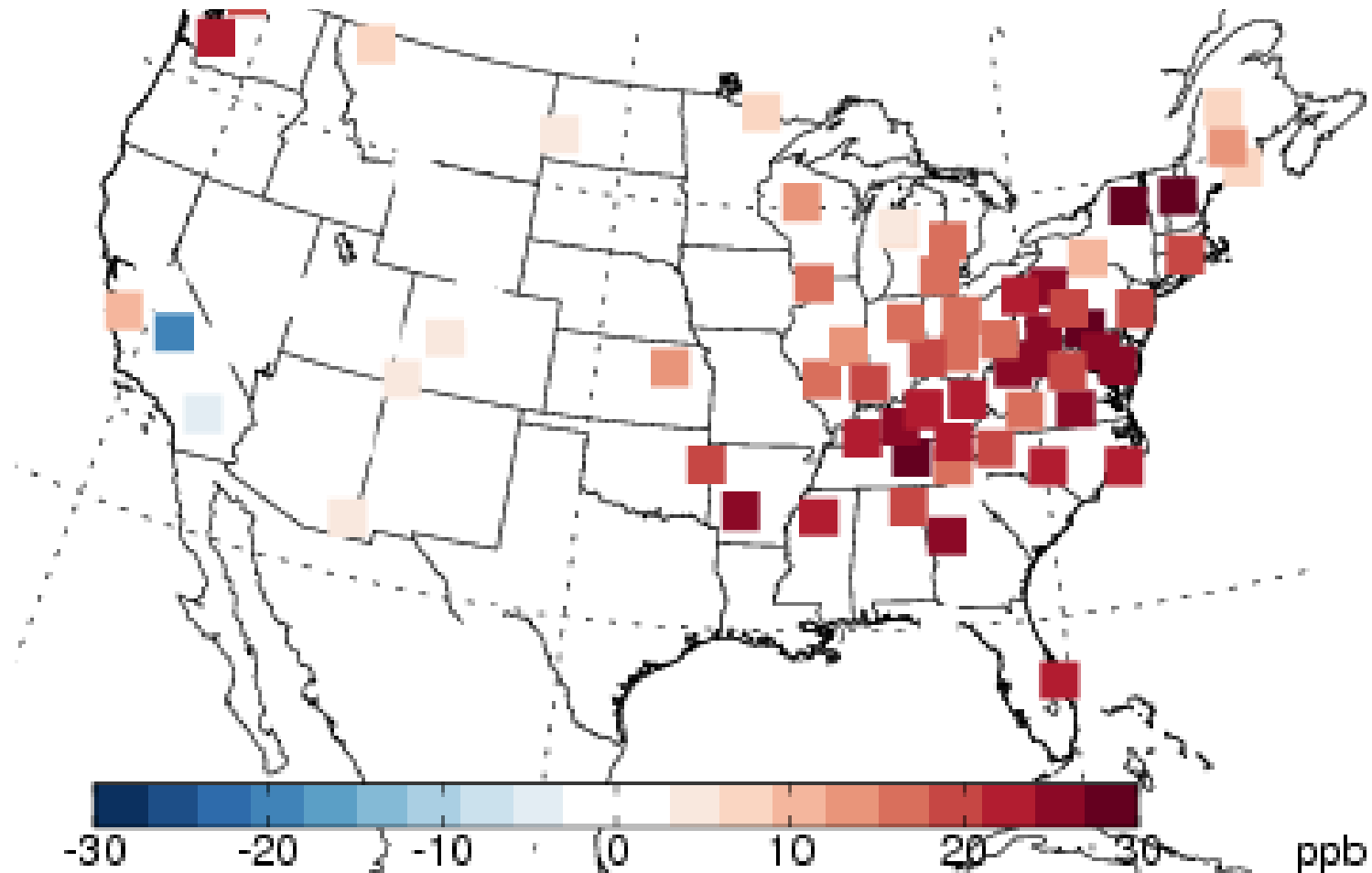
cyclonic ventilation of the eastern U.S.



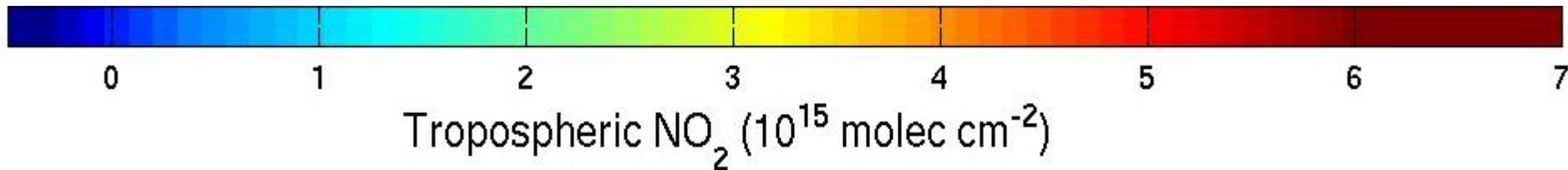
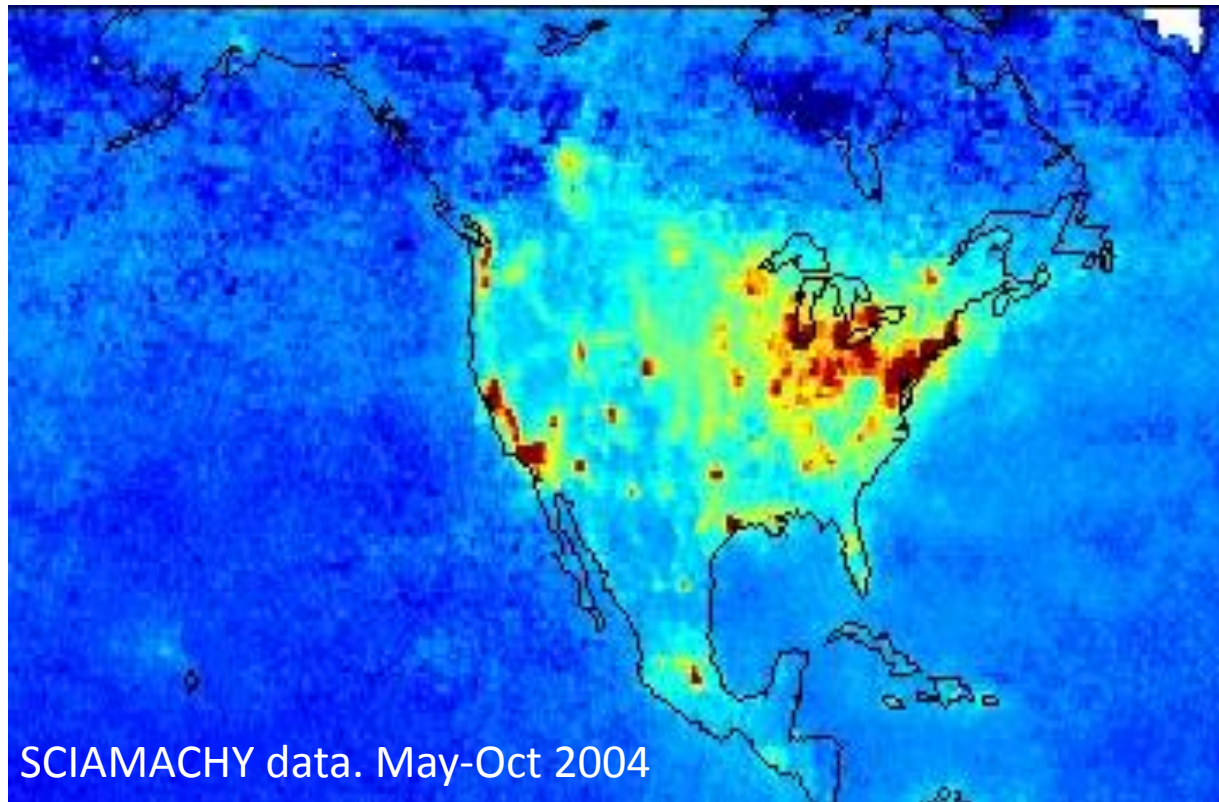
GFDL AM3 CCM eastern U.S. summertime O_3 bias

GFDL AM3 CCM has a high O_3 bias in summer months in the eastern U.S. [10-30 ppb]

June monthly mean MDA8 O_3 Bias (GFDL minus CASTNet)

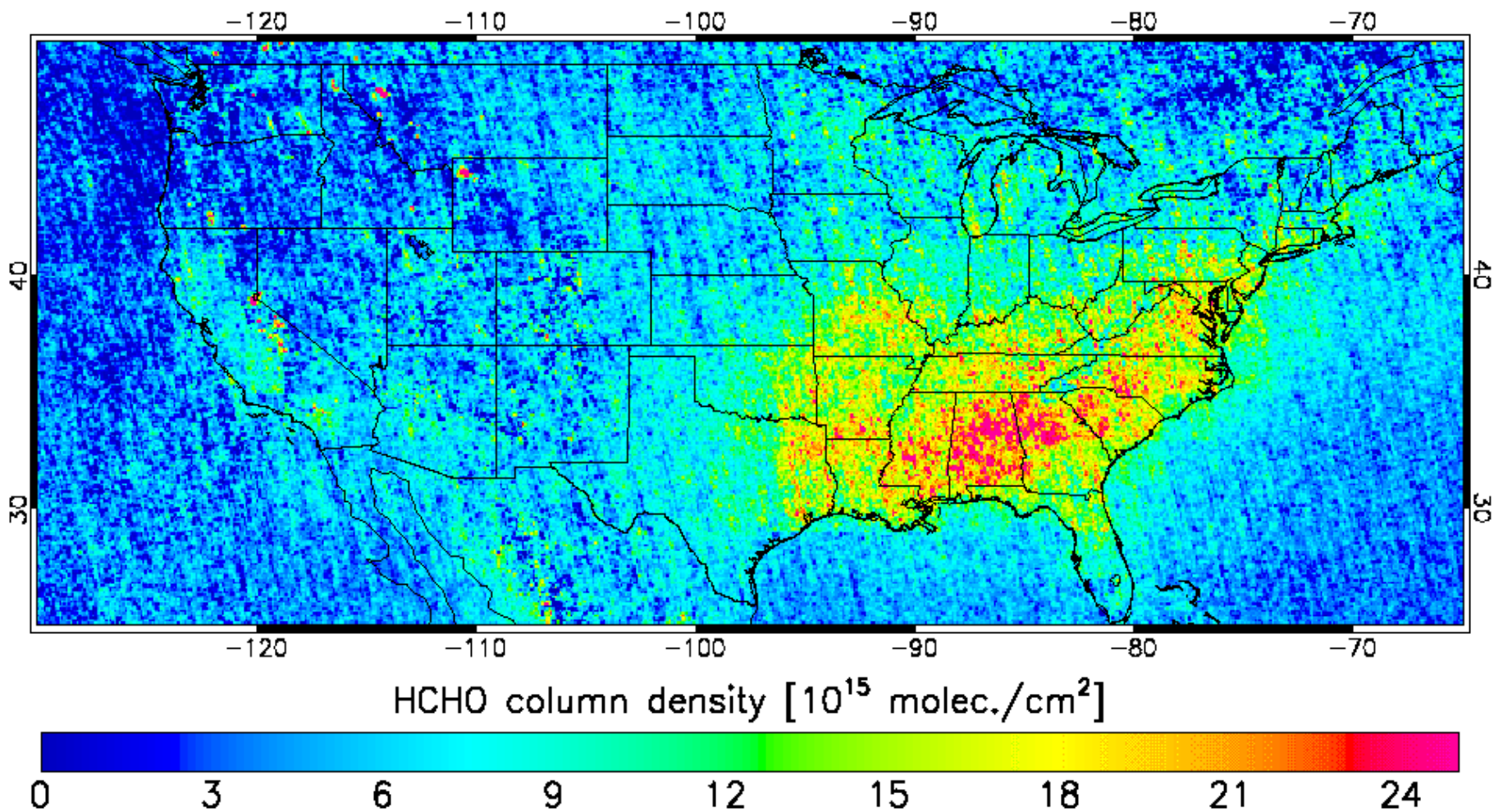


Spatial distribution of NA tropospheric NO₂



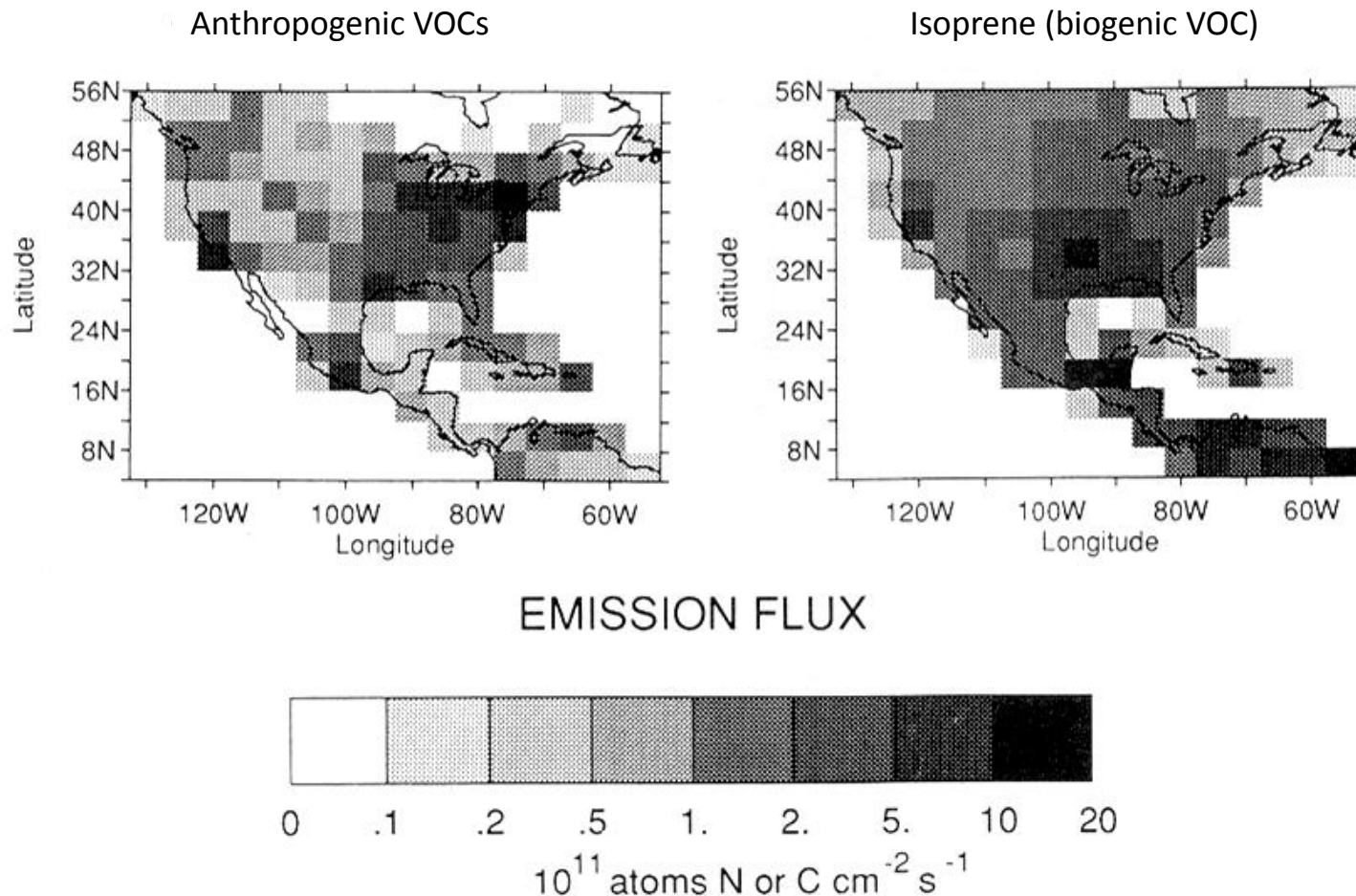
Biogenic VOC (isoprene) column measurements

OMI HCHO: June–August 2006

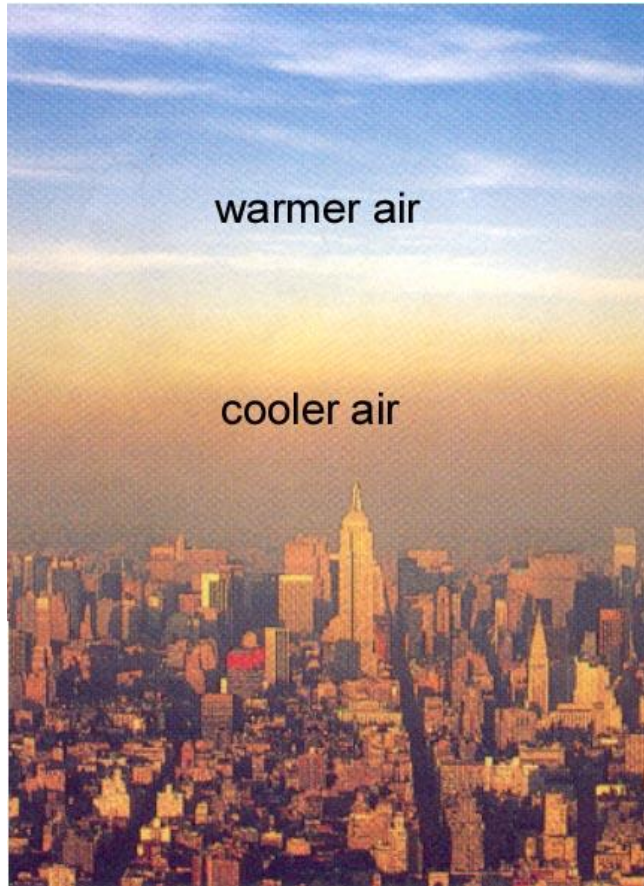


Emissions of VOCs spatially vary

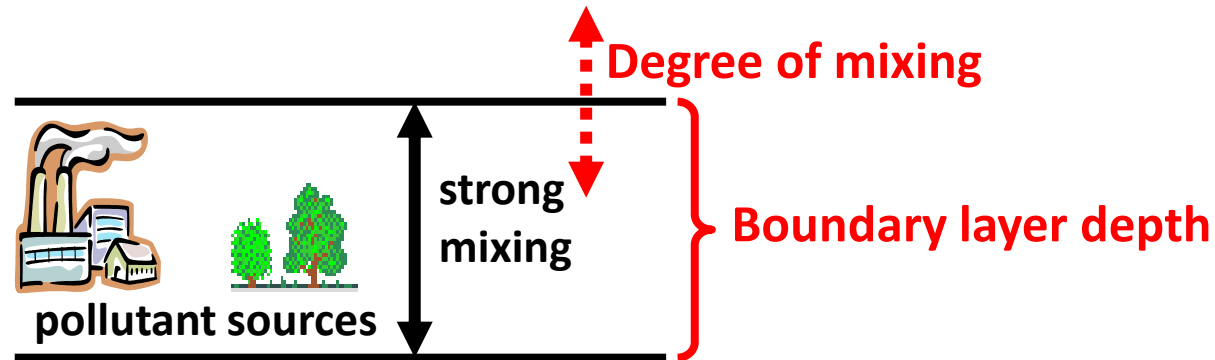
Switches polluted areas in U.S. from NO_x -saturated to NO_x -limited regime!
recognized in Revised Clean Air Act of 1999



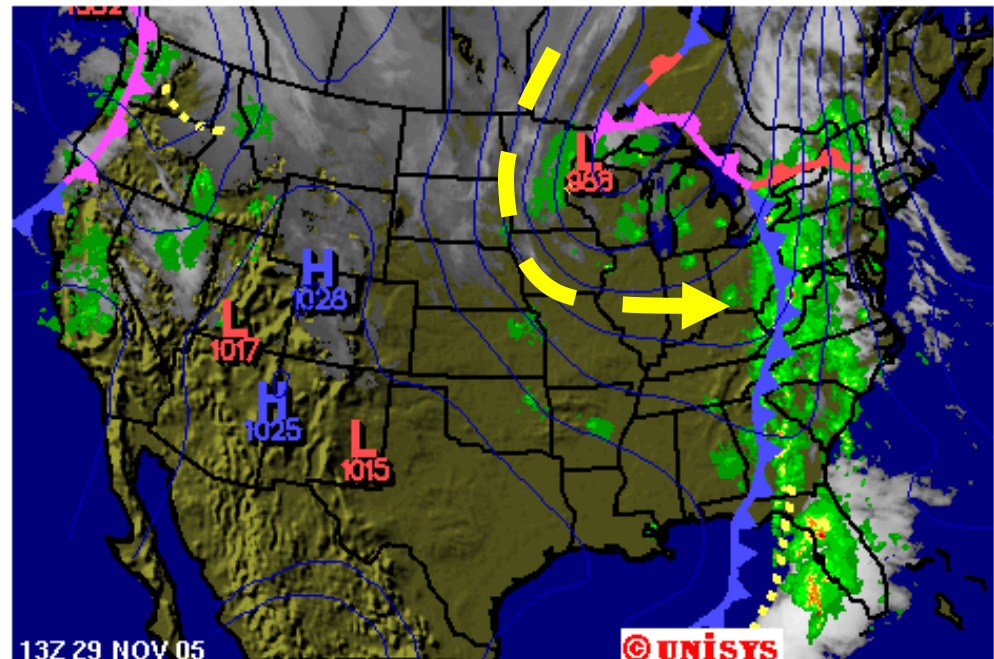
(1) Regional stagnation/ ventilation



[Manhattan, NY]



cyclonic ventilation of the eastern U.S.



(2) PAN chemistry, (3) biogenic emissions

(2) the amount of NO_x ($\text{NO} + \text{NO}_2$) sequestered by peroxyacetyl nitrate (PAN) decreases with increasing temperature

(3) Emissions (biogenic depend strongly on temperature)



Why is O₃ relevant to climate?

Increasing...

Effect on O₃ concentrations:

Stagnation



Temperature



Wind speed



Mixing depth



Cloud cover

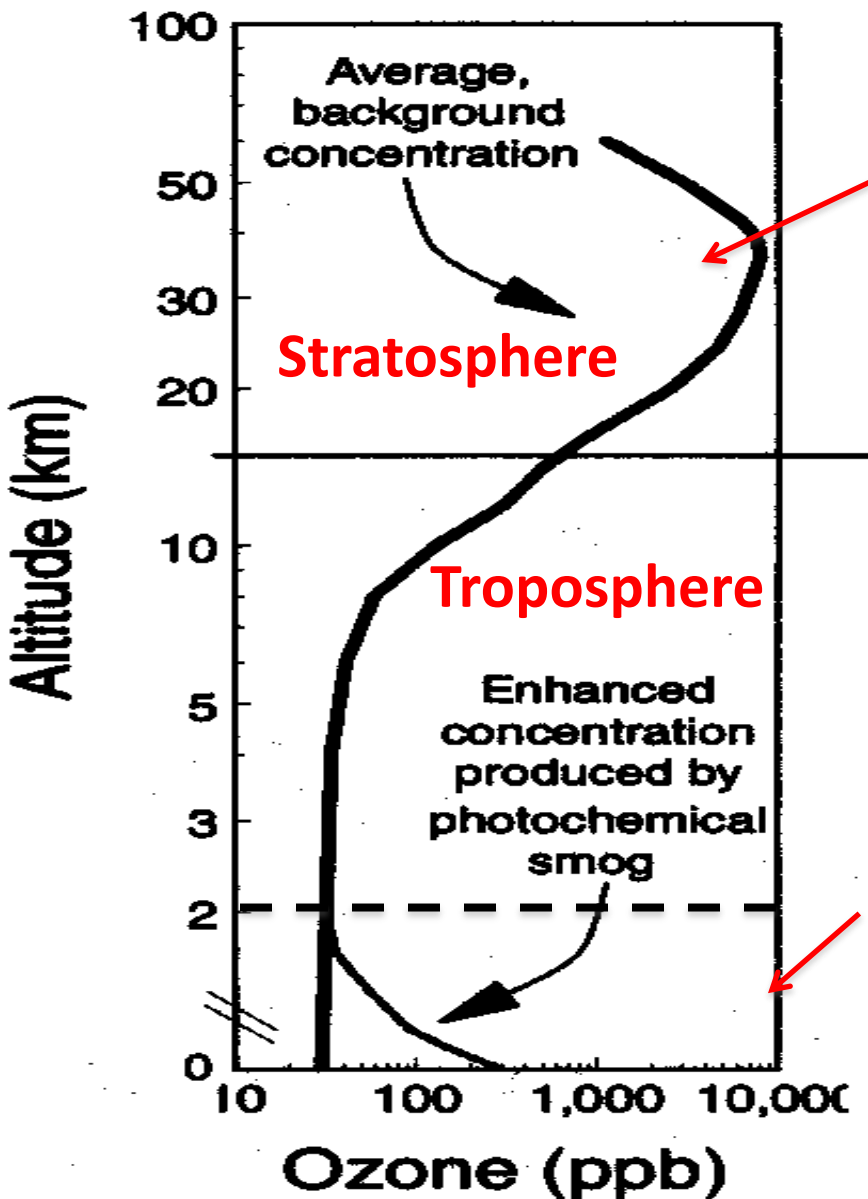


Humidity



[Jacob and Winner, 2009]

Ozone (O_3) in the atmosphere



Good (UV shield) 90% of column O_3 in our atmosphere exists here in the stratosphere

Bad (Smog) surface O_3 is toxic to both people and plants